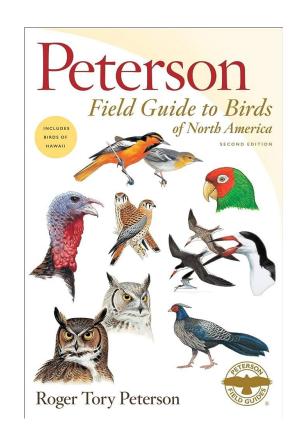


Note: This slide-set is **E**normous—way too much to be fruitfully consumed in a single sesh. That said, it contains numerous natural breaks (ie, section headers), so take advantage of these when you need to. (There's one long stretch without a natural break, so I put a *Break Time* slide within it.)

Many ophthalmologists dread the prospect of interpreting path slides. < raises hand> In this slide-set we will simplify path identification by borrowing from the concept of the birdwatching field guide—reference books that facilitate bird identification by pointing out key characteristics ('field marks') for each species. We will endeavor to do the same for common ophthalmic pathology.



Roger Tory Peterson (1908-1996), creator of the first birding field guide (bottom; top is a Northern Mockingbird)





A page from Peterson's field guide; the lines point to field marks (which are discussed in text on facing pages)



 When you encounter a photomicrograph of the angle, be on the lookout for:

. ?

. ?

Three types of angle issues (not specific conditions)

• 7



The AC Angle



- When you encounter a photomicrograph of the angle, be on the lookout for:
 - Traumatic changes

Dysgeneses

Three types of angle issues (not specific conditions)



- When you encounter a photomicrograph of the angle, be on the lookout for:
 - Traumatic changes
 - ? • ?

Two well-known post-traumatic angle issues

Dysgeneses

- 7
- When you encounter a photomicrograph of the angle, be on the lookout for:
 - Traumatic changes
 - Cyclodialysis
 - Angle recession

Two well-known post-traumatic angle issues

Dysgeneses



- When you encounter a photomicrograph of the angle, be on the lookout for:
 - Traumatic changes
 - Cyclodialysis
 - Angle recession
 - Dysgeneses
 - . ?
 - . ?

Two well-known dysgeneses



- When you encounter a photomicrograph of the angle, be on the lookout for:
 - Traumatic changes
 - Cyclodialysis
 - Angle recession
 - Dysgeneses
 - Peters anomalyAxenfeld-Rieger

Two well-known dysgeneses



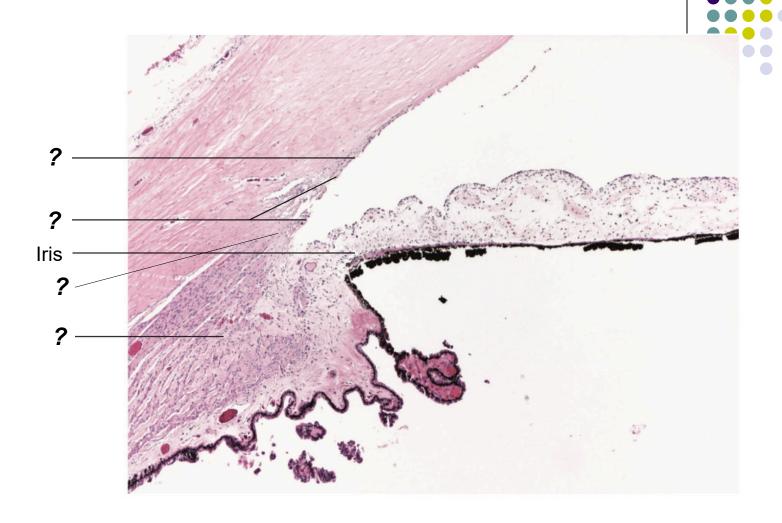
- When you encounter a photomicrograph of the angle, be on the lookout for:
 - Traumatic changes
 - Cyclodialysis
 - Angle recession
 - Dysgeneses
 - Peters anomaly
 - Axenfeld-Rieger
 - Cell clogging the TM
 - ?
 - ?

Two cell types notorious for clogging the TM



- When you encounter a photomicrograph of the angle, be on the lookout for:
 - Traumatic changes
 - Cyclodialysis
 - Angle recession
 - Dysgeneses
 - Peters anomaly
 - Axenfeld-Rieger
 - Cell clogging the TM
 - RBCs
 - Macrophages

Two cell types notorious for clogging the TM

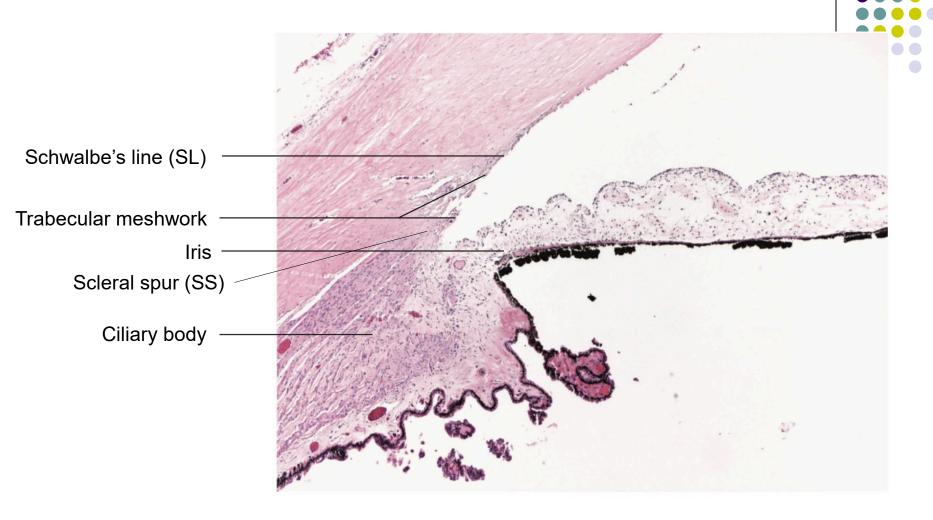


12

Question

These are the key angle landmarks—name them. (I assume you've got the iris on lock already.)

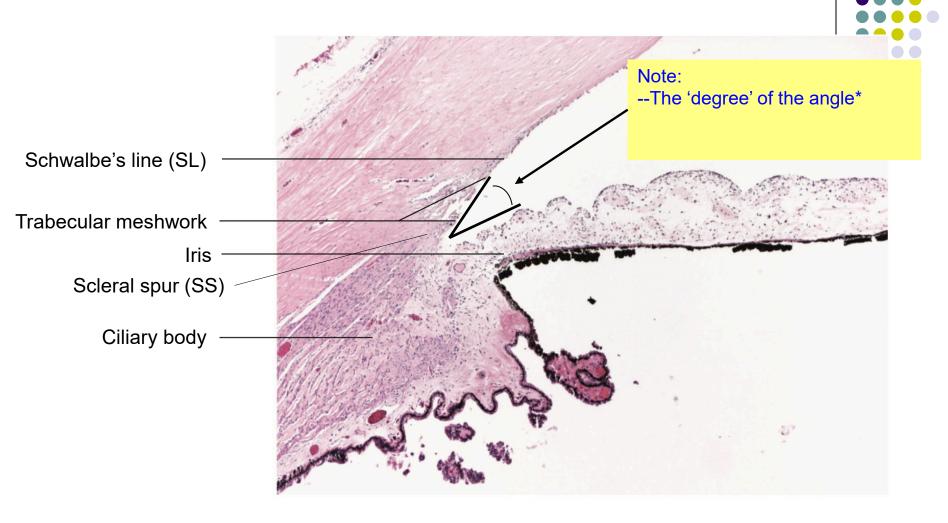
13





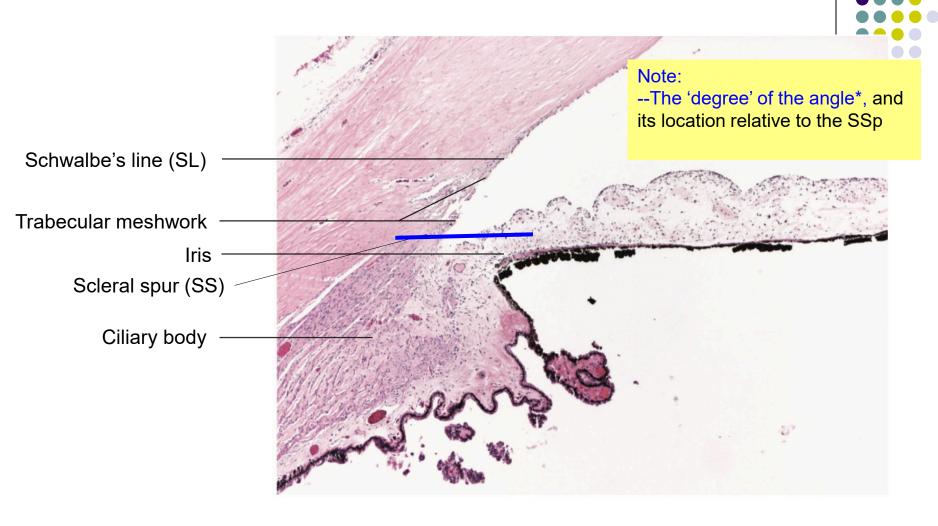
These are the key angle landmarks—name them. (I assume you've got the iris on lock already.)

14



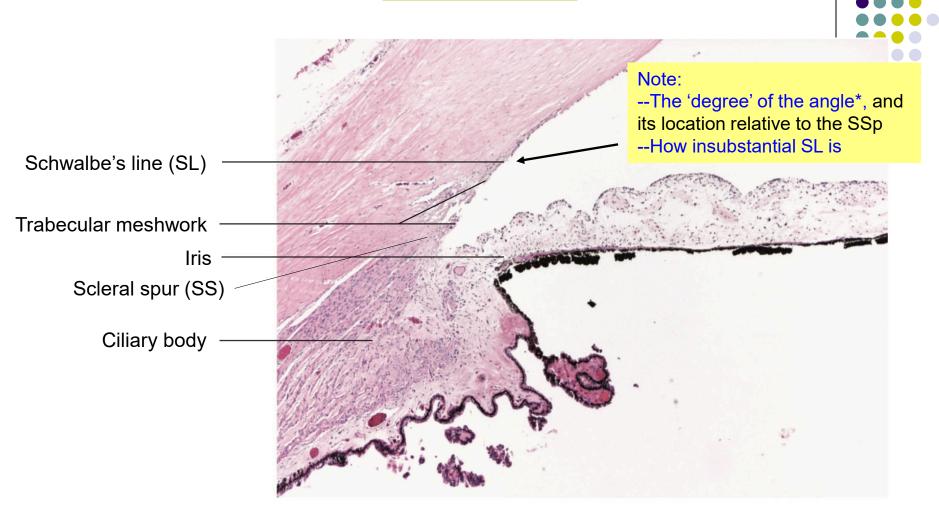
These are the key angle landmarks—name them. (I assume you've got the iris on lock already.)

*Bearing in mind there is considerable anatomic variability in the degree of the normal angle



These are the key angle landmarks—name them. (I assume you've got the iris on lock already.)

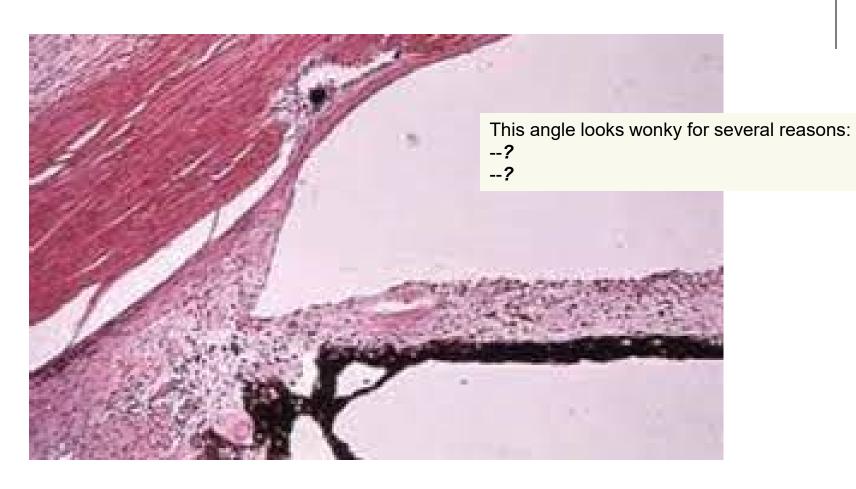
15



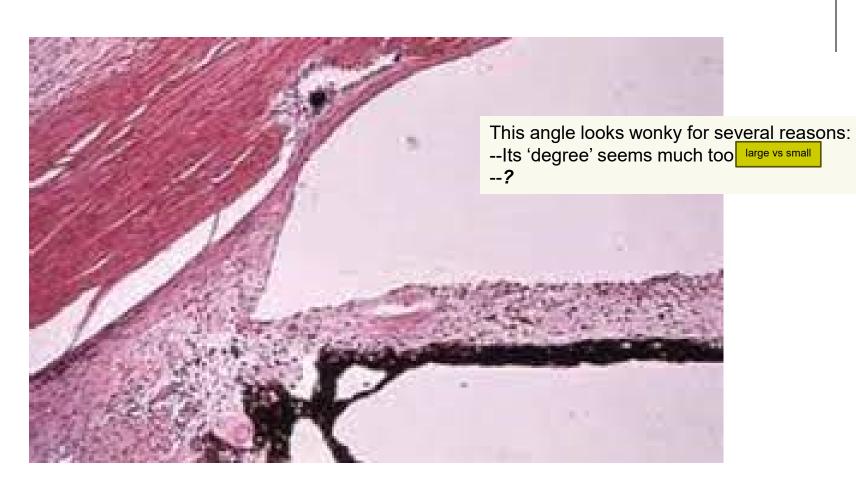
These are the key angle landmarks—name them. (I assume you've got the iris on lock already.)

16

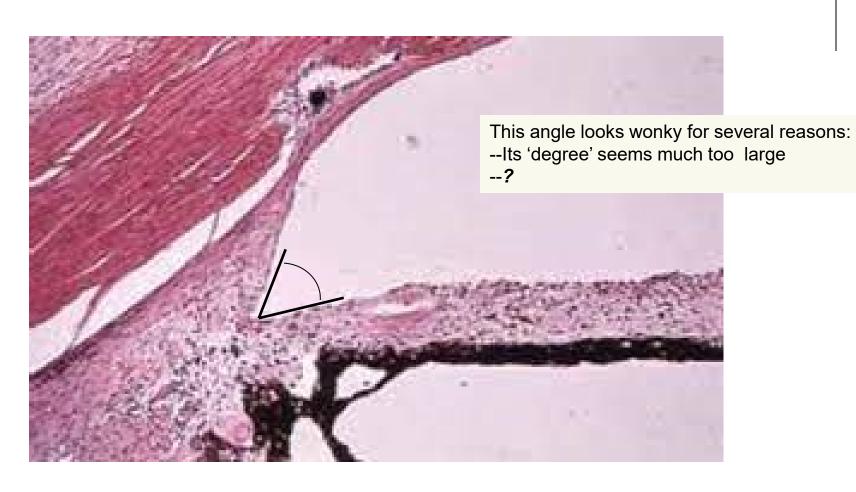




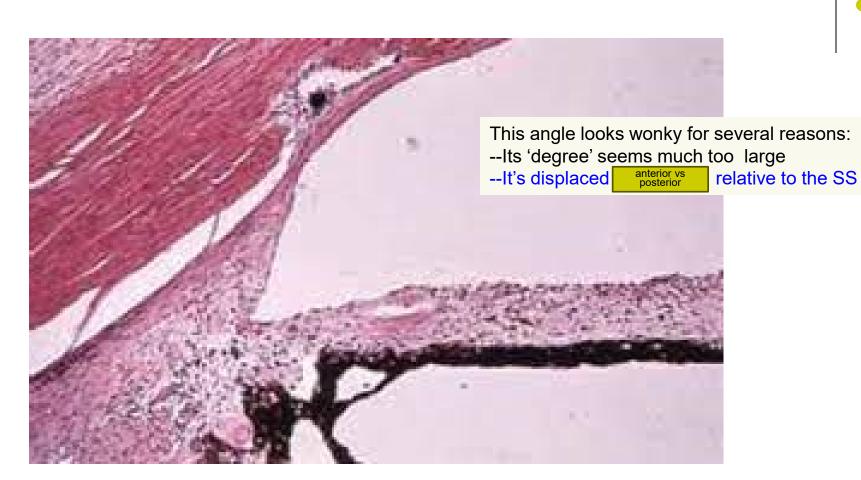




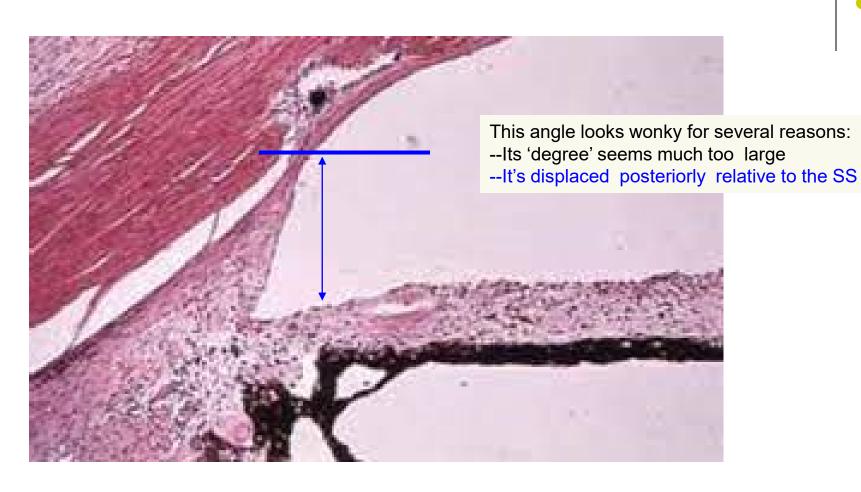


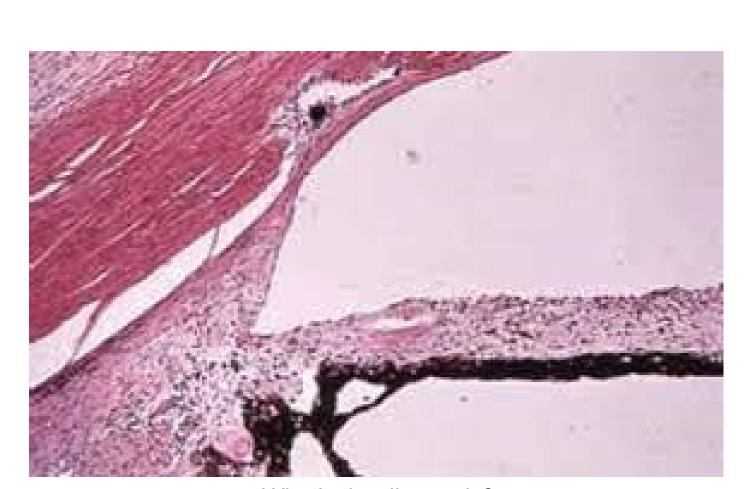








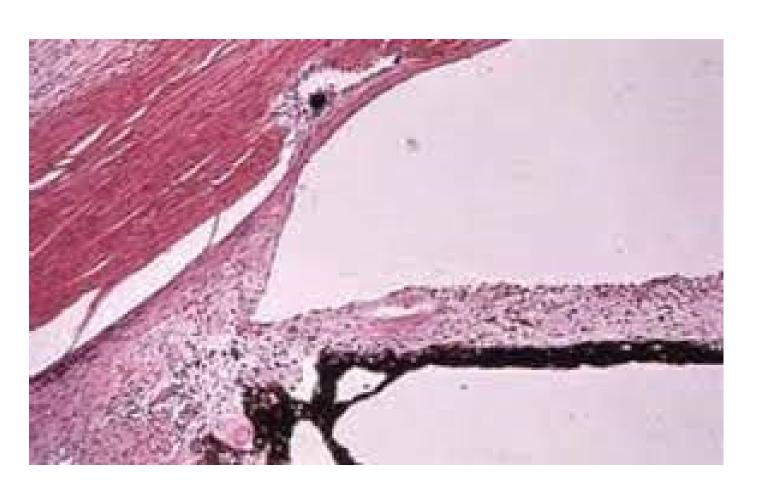




What's the diagnosis?



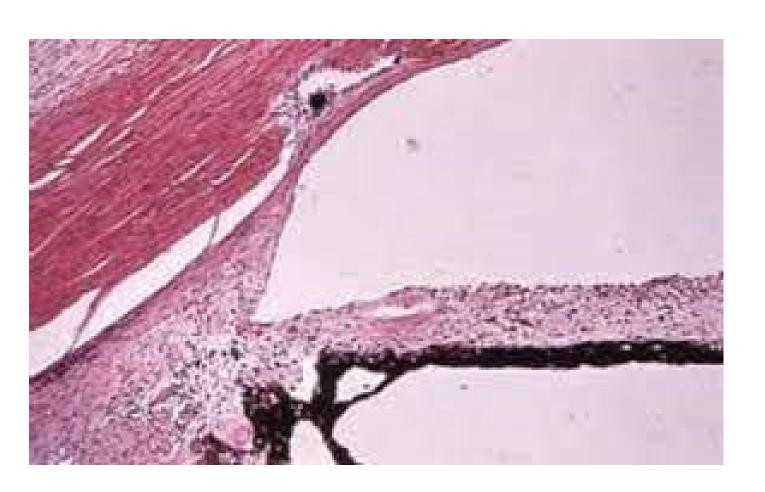




Angle recession. Blunt trauma has torn the

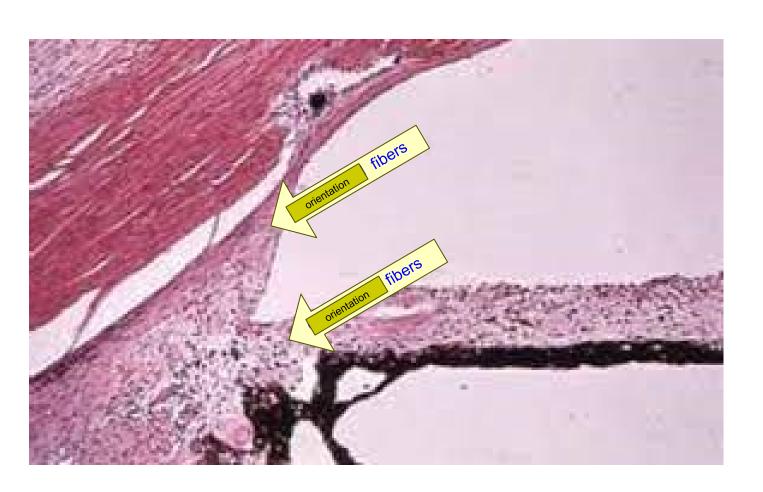
structure





Angle recession. Blunt trauma has torn the ciliary body (CB)

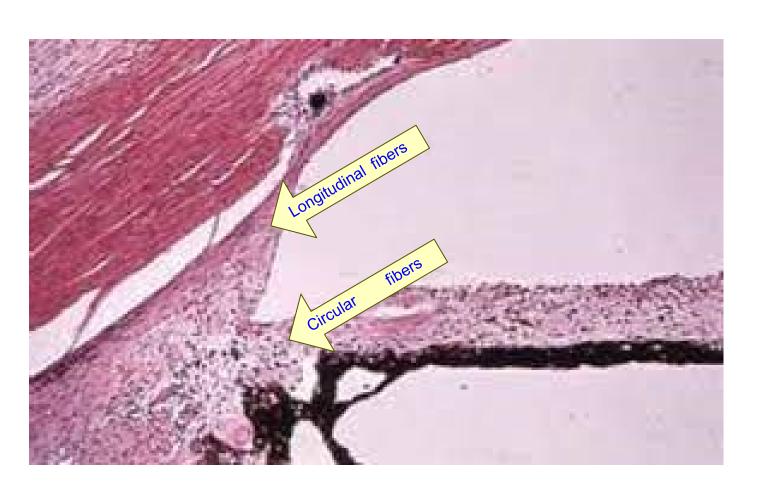




Angle recession. Blunt trauma has torn the ciliary body (CB), tearing its fibers away from its orientation fibers.

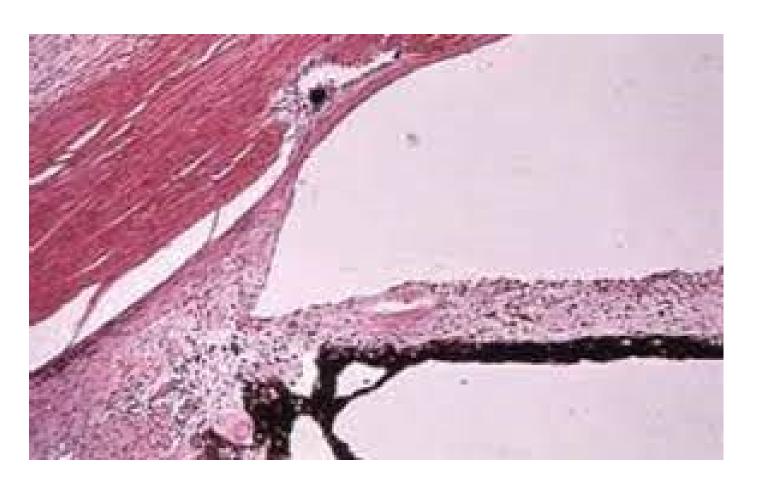
orientation





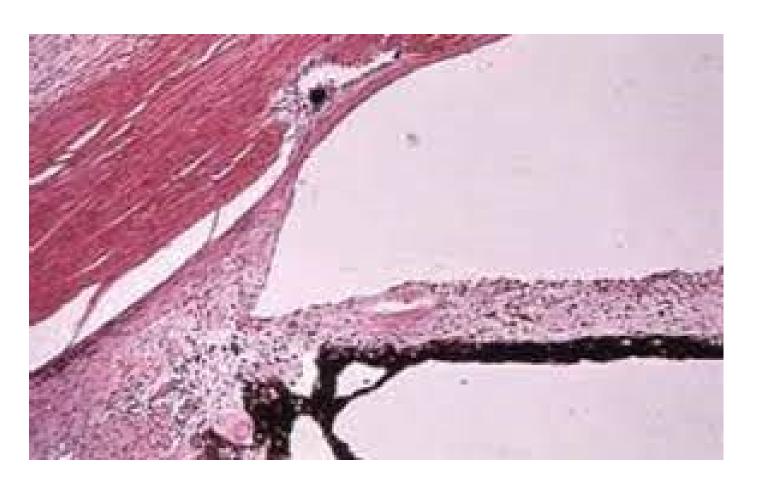
Angle recession. Blunt trauma has torn the ciliary body (CB) , tearing its longitudinal fibers away from its circular fibers.





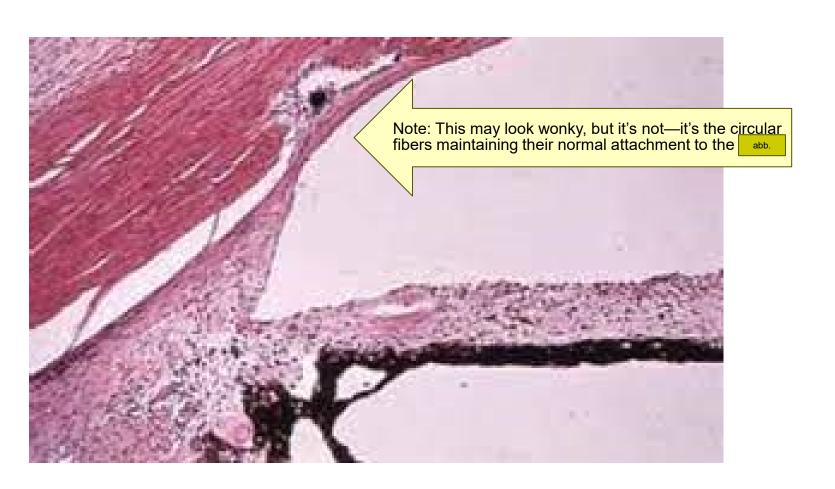
Angle recession. Blunt trauma has torn the ciliary body (CB), tearing its longitudinal fibers away from its circular fibers. Such pts are at high risk for developing





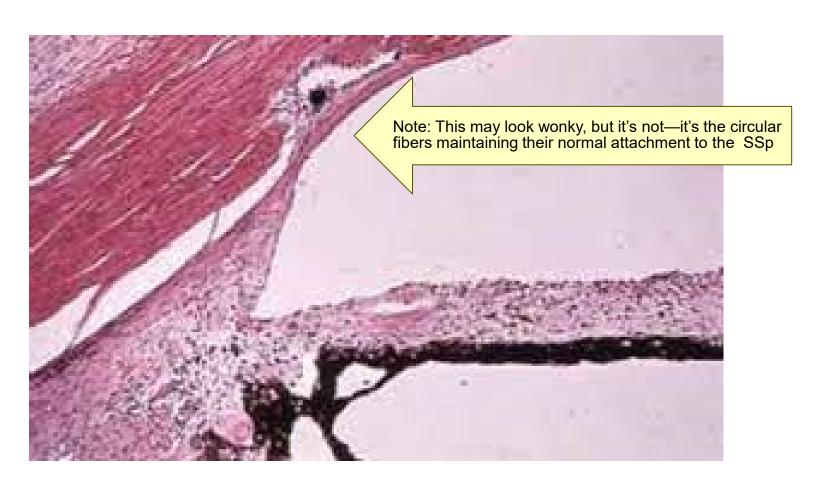
Angle recession. Blunt trauma has torn the ciliary body (CB), tearing its longitudinal fibers away from its circular fibers. Such pts are at high risk for developing glaucoma





Angle recession. Blunt trauma has torn the ciliary body (CB), tearing its longitudinal fibers away from its circular fibers. Such pts are at high risk for developing glaucoma



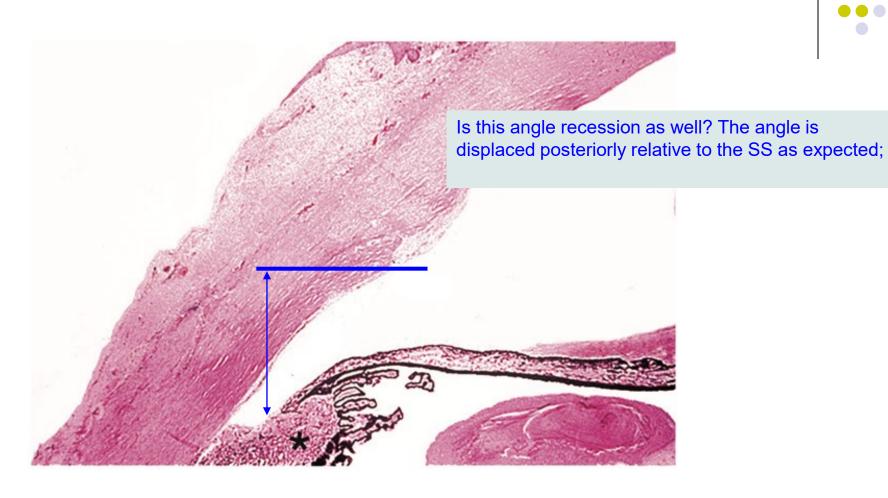


Angle recession. Blunt trauma has torn the ciliary body (CB), tearing its longitudinal fibers away from its circular fibers. Such pts are at high risk for developing glaucoma

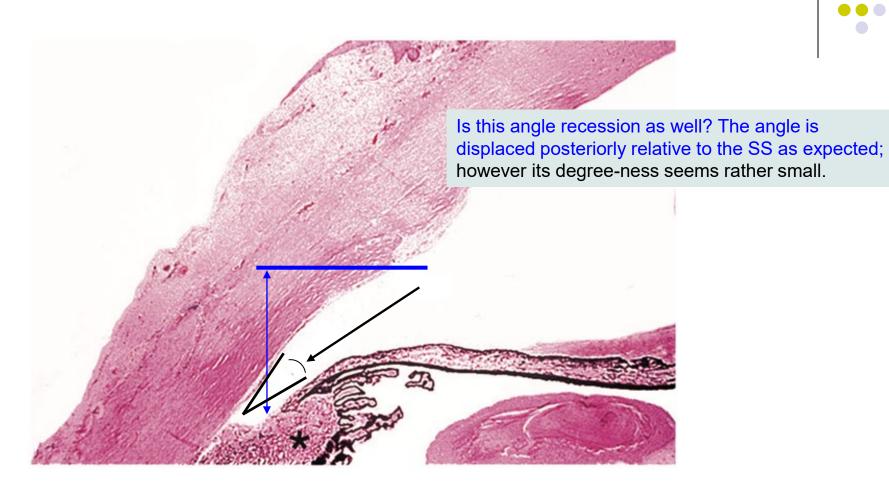


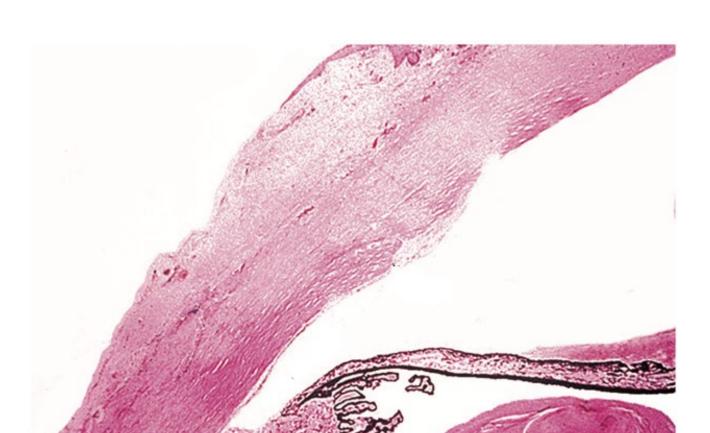










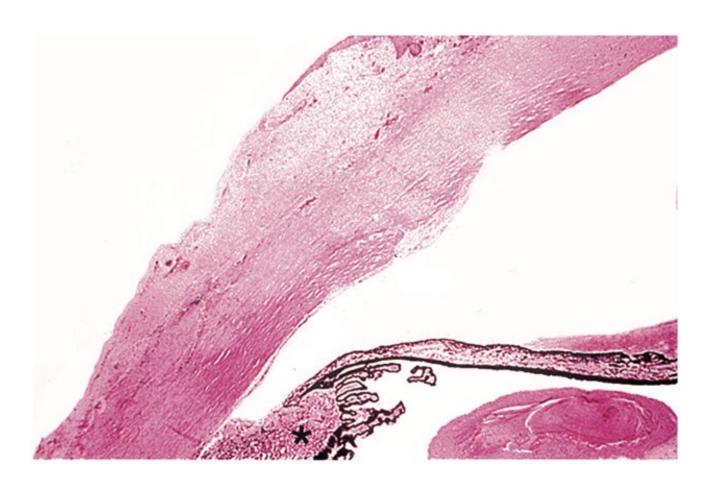


What's the diagnosis?



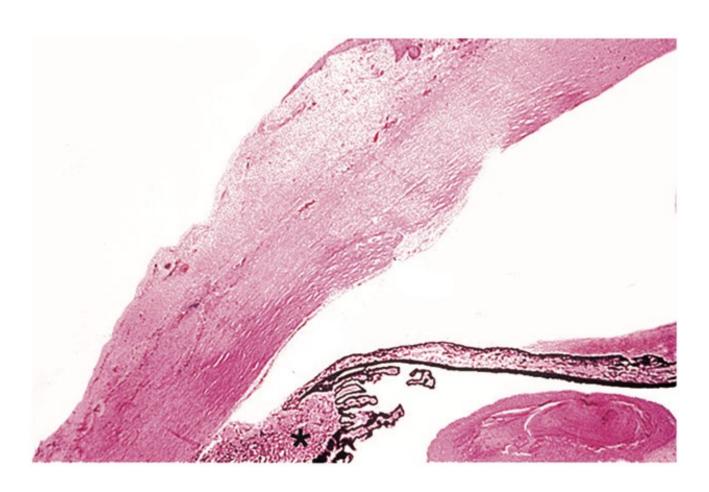


35



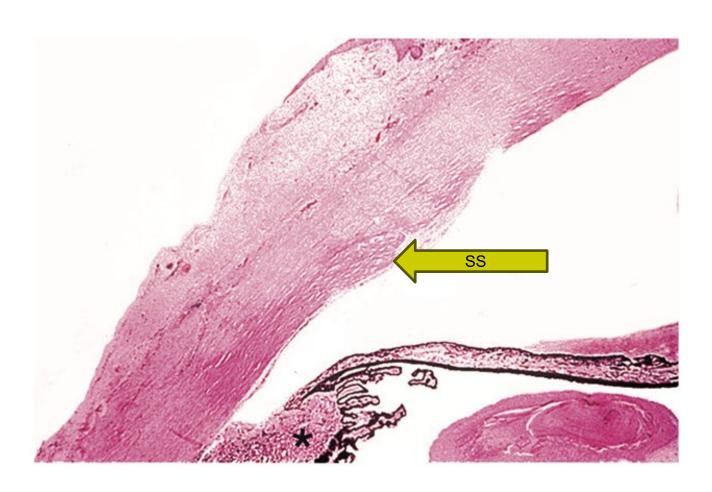
Cyclodialysis. Blunt trauma has again torn the ciliary body (asterisk)





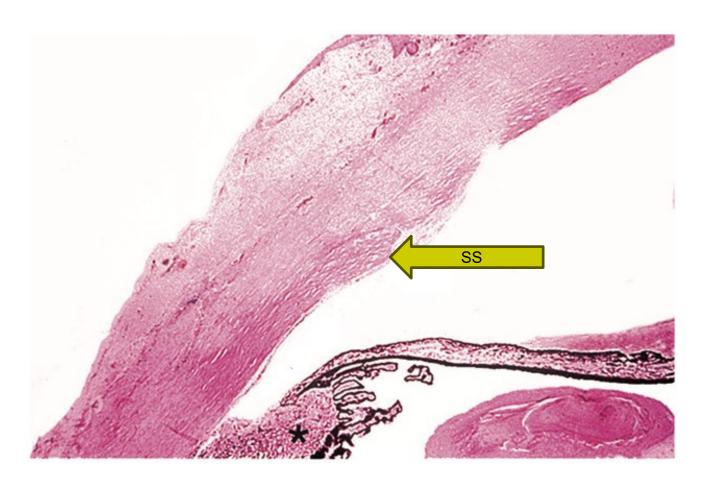
Cyclodialysis. Blunt trauma has again torn the ciliary body (asterisk), but rather than tearing fiber from fiber a la recession, it has torn away from its normal attachment to the labb.





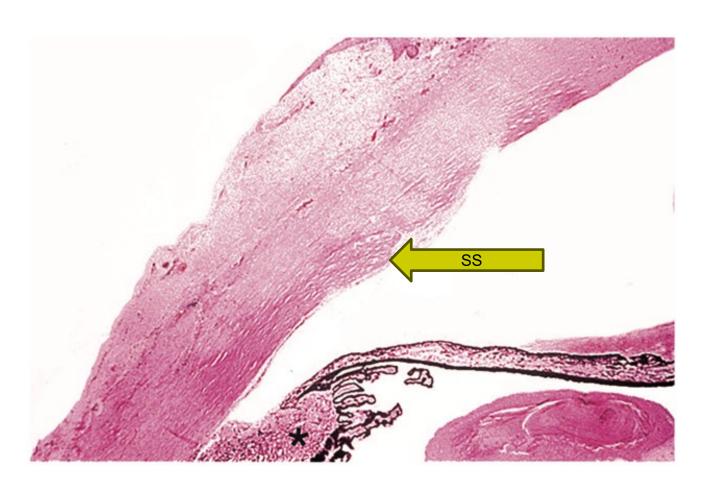
Cyclodialysis. Blunt trauma has again torn the ciliary body (asterisk), but rather than tearing fiber from fiber a la recession, it has torn away from its normal attachment to the SS





Cyclodialysis. Blunt trauma has again torn the ciliary body (asterisk), but rather than tearing fiber from fiber a la recession, it has torn away from its normal attachment to the SS. Such pts are also at risk for developing glaucoma (less v more) so than those with angle recession).





Cyclodialysis. Blunt trauma has again torn the ciliary body (asterisk), but rather than tearing fiber from fiber a la recession, it has torn away from its normal attachment to the SS. Such pts are also at risk for developing glaucoma (less so than those with angle recession).





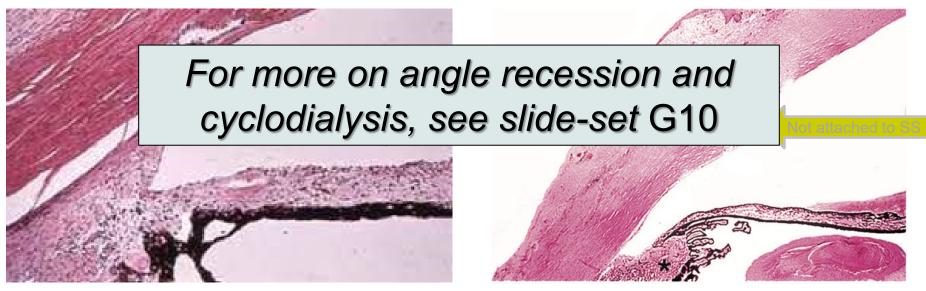


Angle recession

Cyclodialysis

Angle recession and cyclodialysis side-by-side



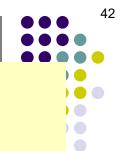


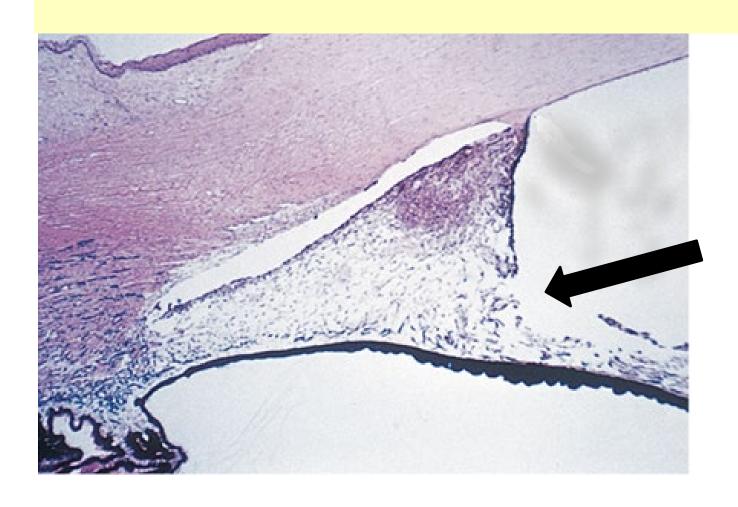
Angle recession

Cyclodialysis

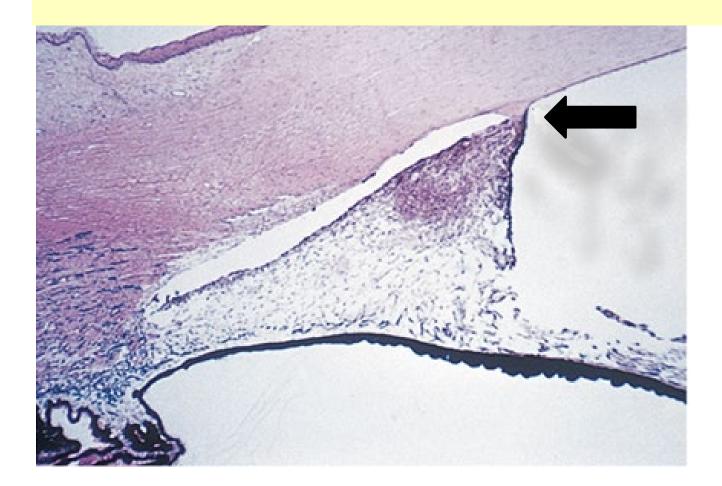
Angle recession and cyclodialysis side-by-side

At first glance, this looks like angle recession, with what appears to be a split in the CB



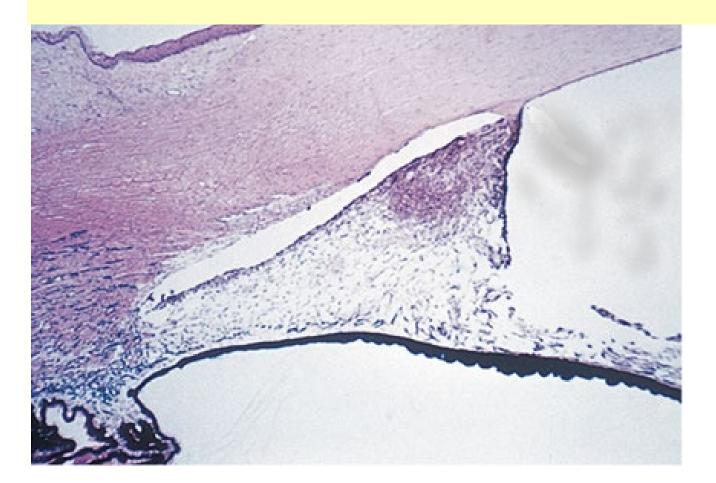


At first glance, this looks like angle recession, with what appears to be a split in the CB and an intact CB-SS attachment.



At first glance, this looks like angle recession, with what appears to be a split in the CB and an intact CB-SS attachment. But careful inspection of the image reveals problems with this interpretation:

--The TM is [locate it]

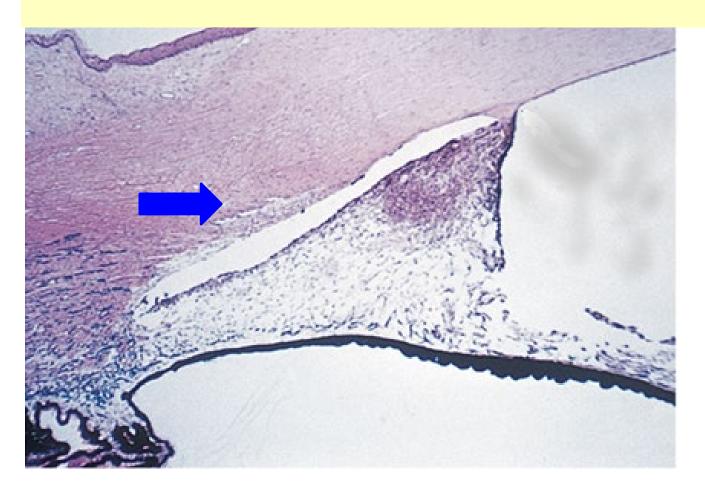


At first glance, this looks like angle recession, with what appears to be a split in the CB and an intact CB-SS attachment. But careful inspection of the image reveals problems with this interpretation:



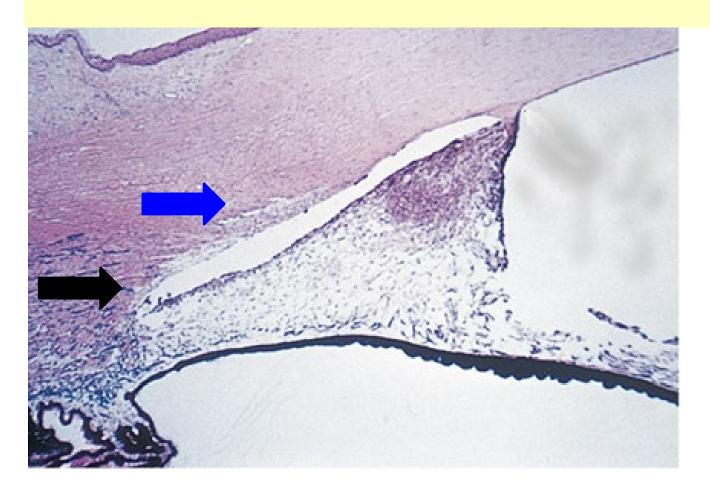
At first glance, this looks like angle recession, with what appears to be a split in the CB and an intact CB-SS attachment. But careful inspection of the image reveals problems with this interpretation:

--The TM is here, which means the SS is about [ditto]

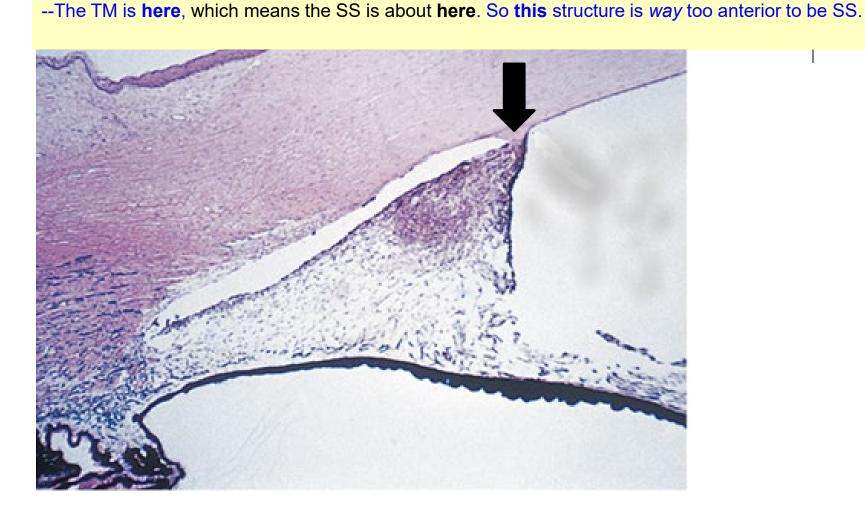


At first glance, this looks like angle recession, with what appears to be a split in the CB and an intact CB-SS attachment. But careful inspection of the image reveals problems with this interpretation:

--The TM is here, which means the SS is about here.



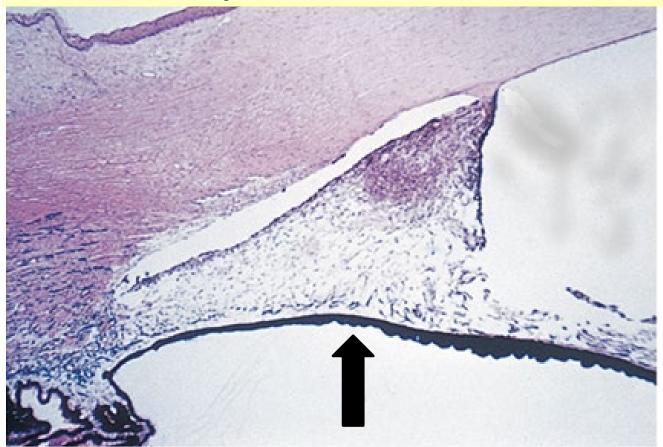
At first glance, this looks like angle recession, with what appears to be a split in the CB and an intact CB-SS attachment. But careful inspection of the image reveals problems with this interpretation:

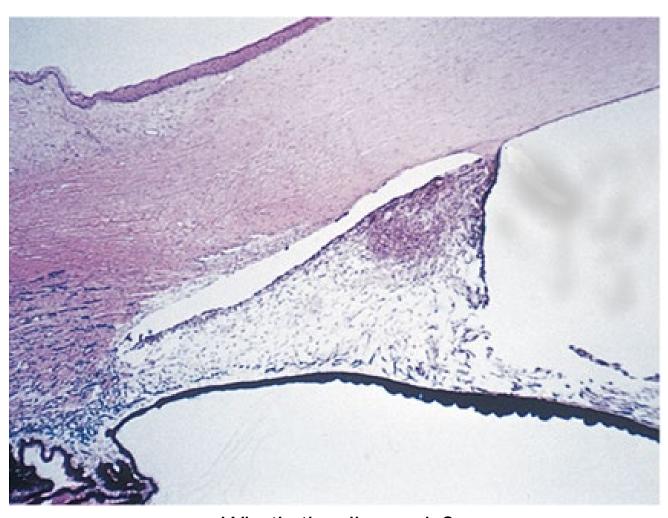




At first glance, this looks like angle recession, with what appears to be a split in the CB and an intact CB-SS attachment. But careful inspection of the image reveals problems with this interpretation:

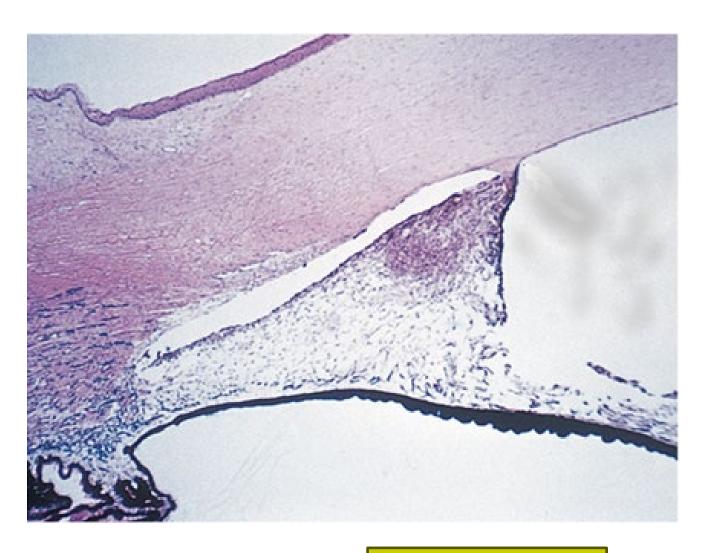
- --The TM is here, which means the SS is about here. So this structure is way too anterior to be SS.
- --The attachment is arising from the *iris*, not the CB.





What's the diagnosis?

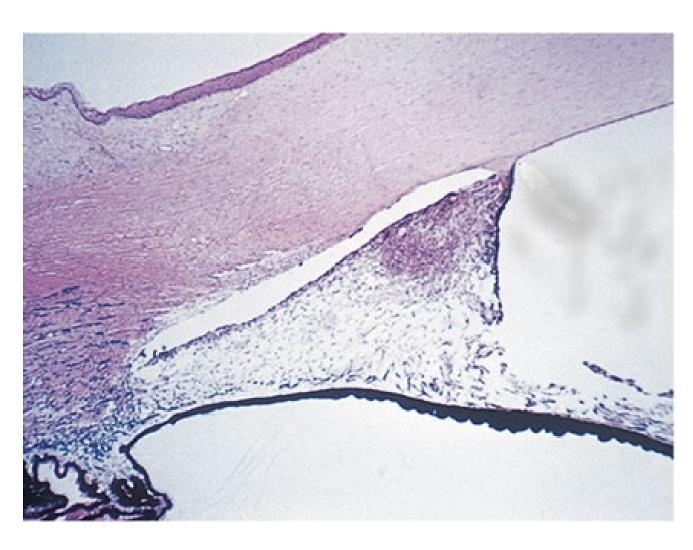






class of condition (three words)

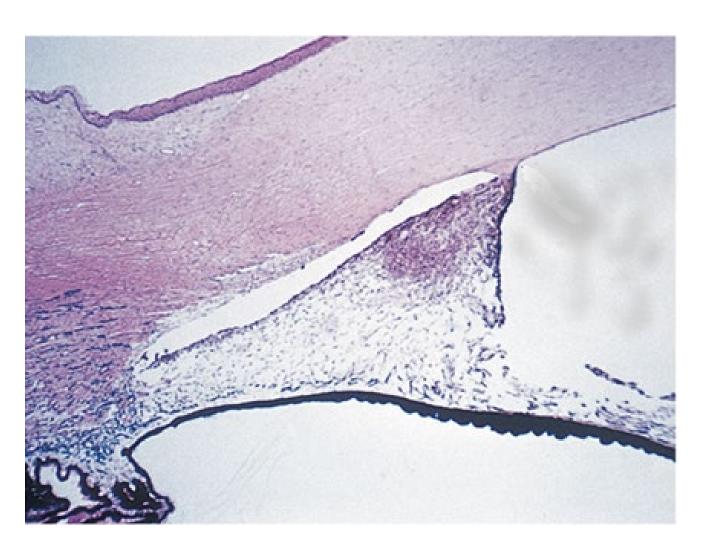




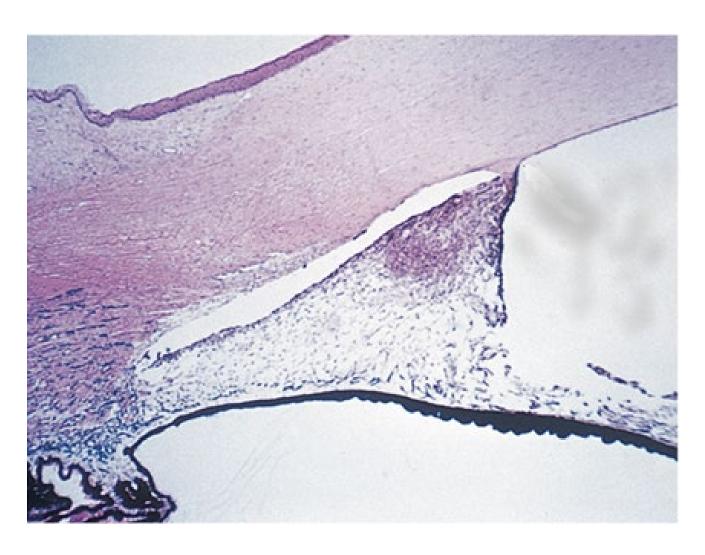
Axenfeld-Rieger syndrome, an anterior-segment dysgenesis.





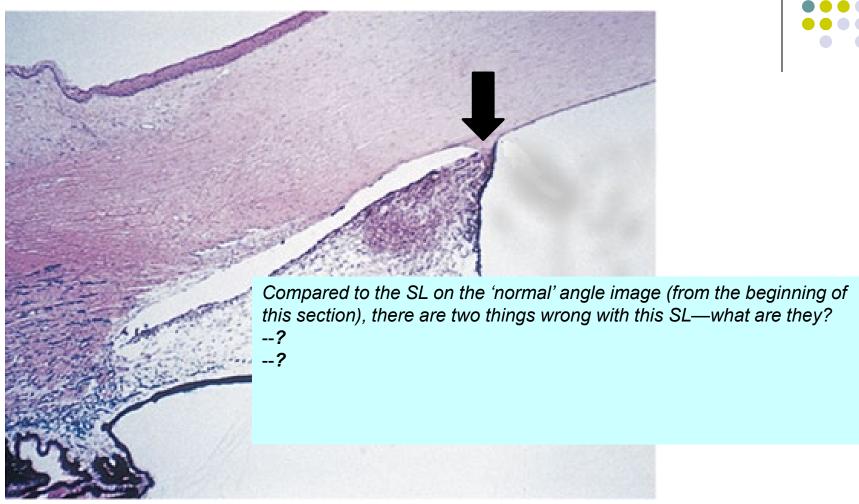




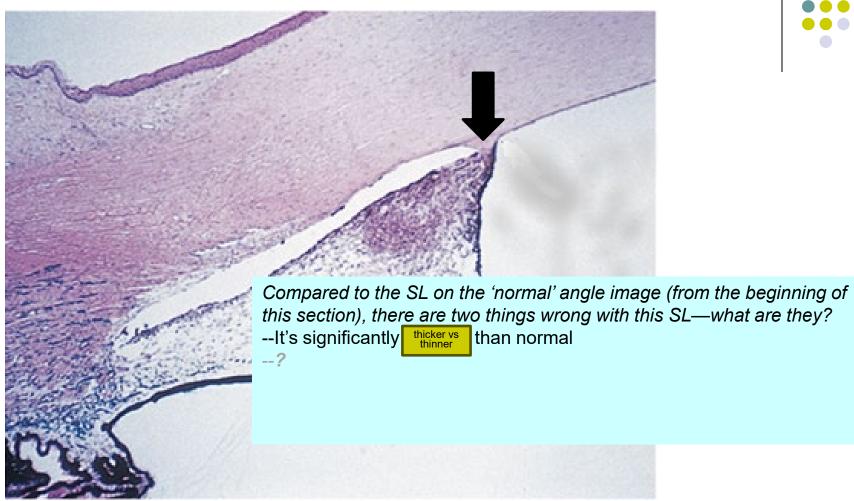


Axenfeld-Rieger syndrome, an anterior-segment dysgenesis. Rather than CB fibers attaching to the SS, the attachment is an iris process attaching to SL.

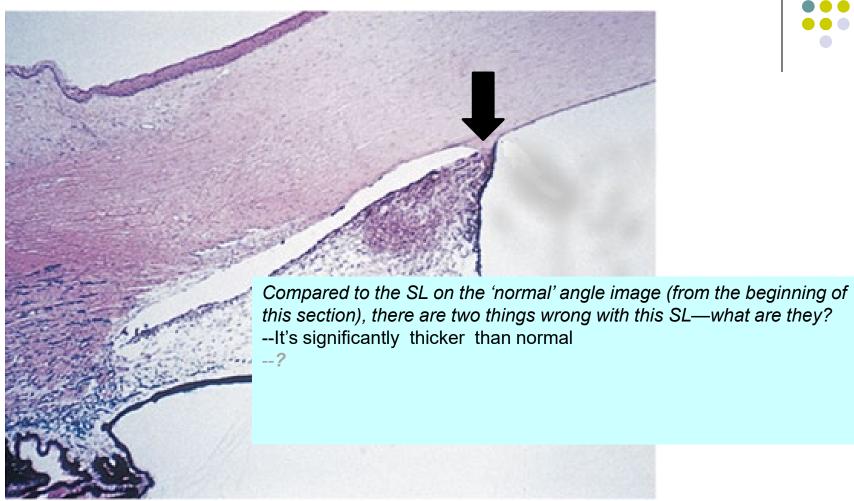




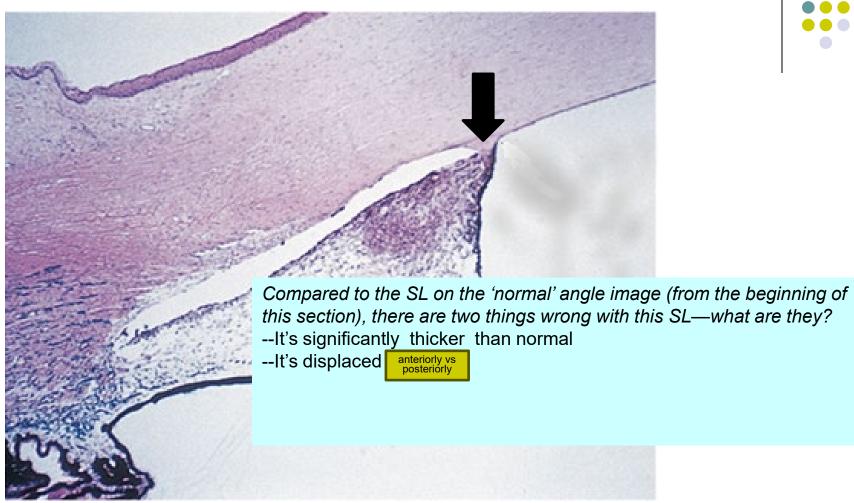




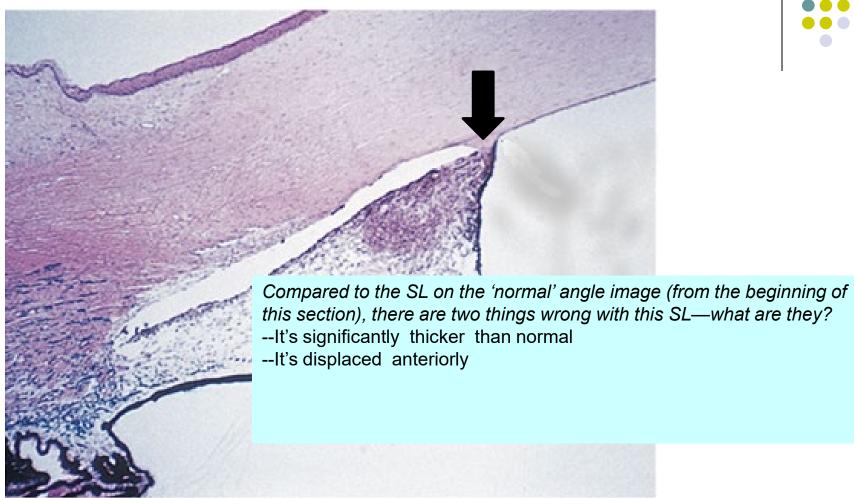




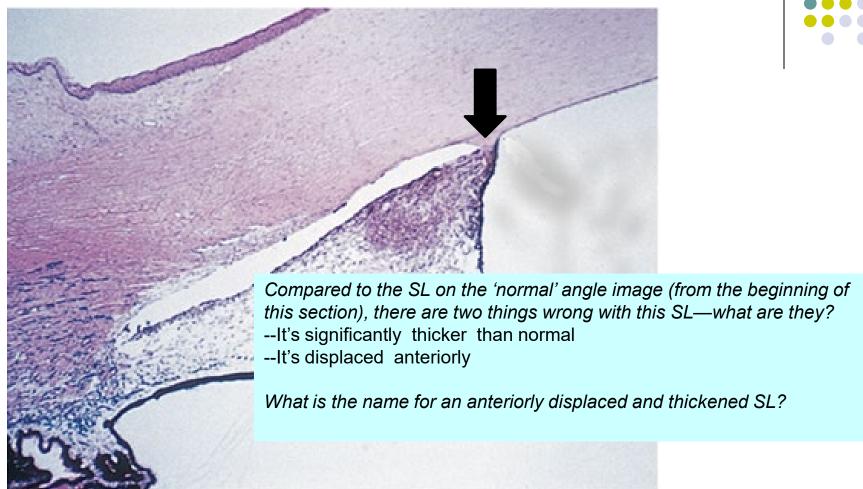




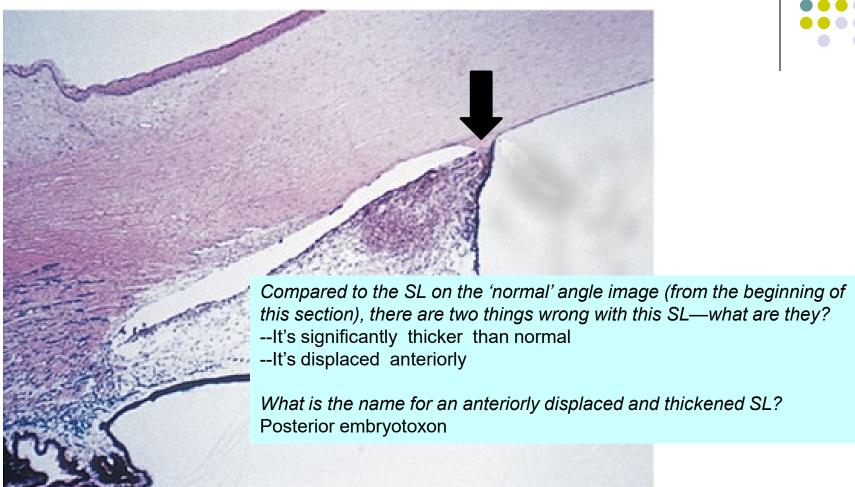






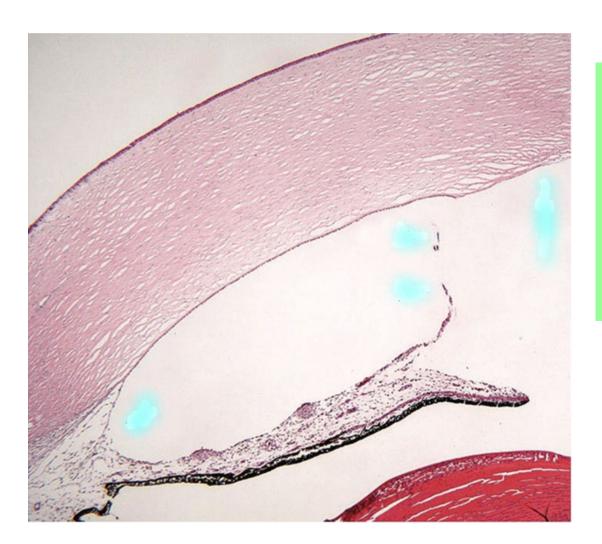






62

Pathwatching





At first glance this looks like Axenfeld-Rieger as well, with what appears to be an iris process attaching to a posterior embryotoxon. But close inspection reveals problems with this:

--?

--?

63

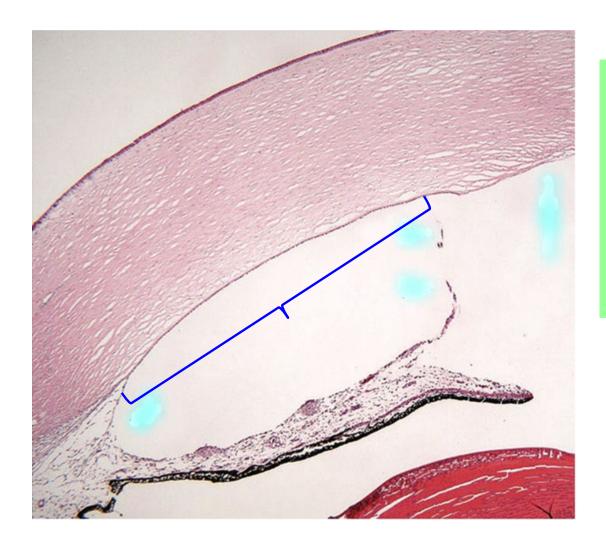
Pathwatching





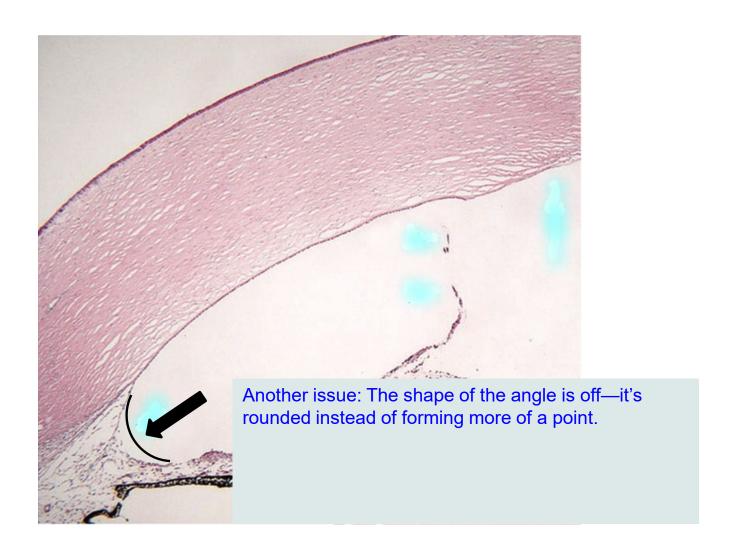
At first glance this looks like Axenfeld-Rieger as well, with what appears to be an iris process attaching to a posterior embryotoxon. But close inspection reveals problems with this:

--A 'SL' doesn't seem to be present at all (much less a thickened one).

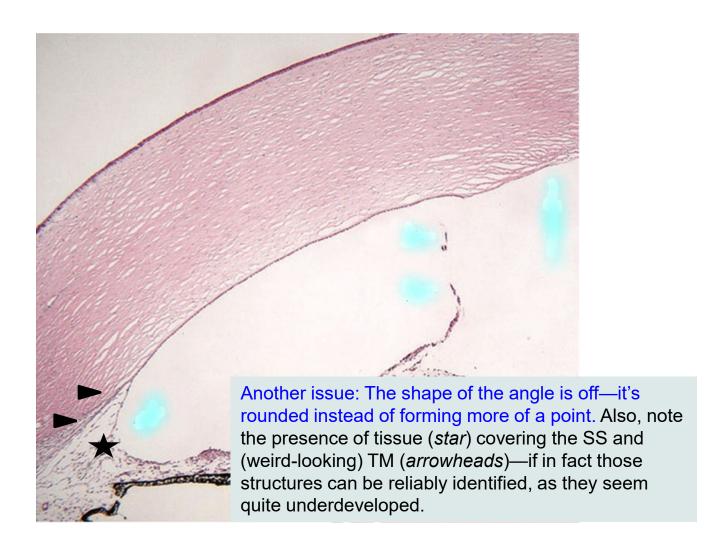




At first glance this looks like
Axenfeld-Rieger as well, with what
appears to be an iris process
attaching to a posterior
embryotoxon. But close inspection
reveals problems with this:
--A 'SL' doesn't seem to be present
at all (much less a thickened one).
--The point of attachment is way
more anterior than we saw on the
A-R slide.

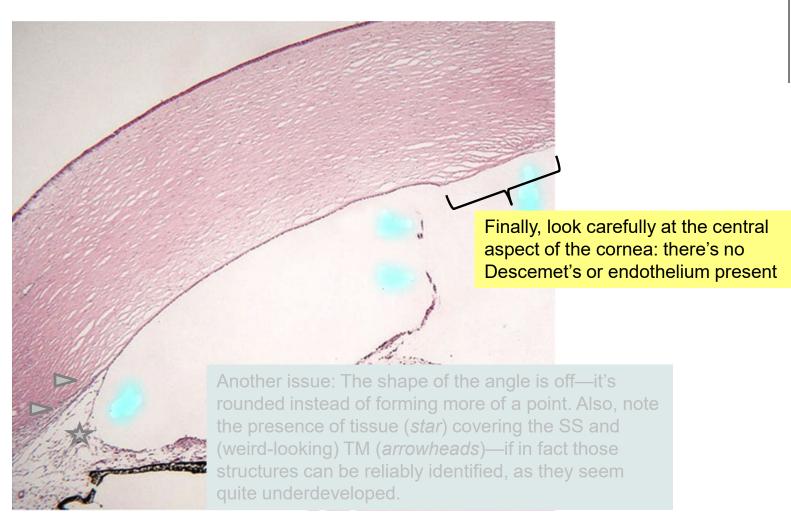




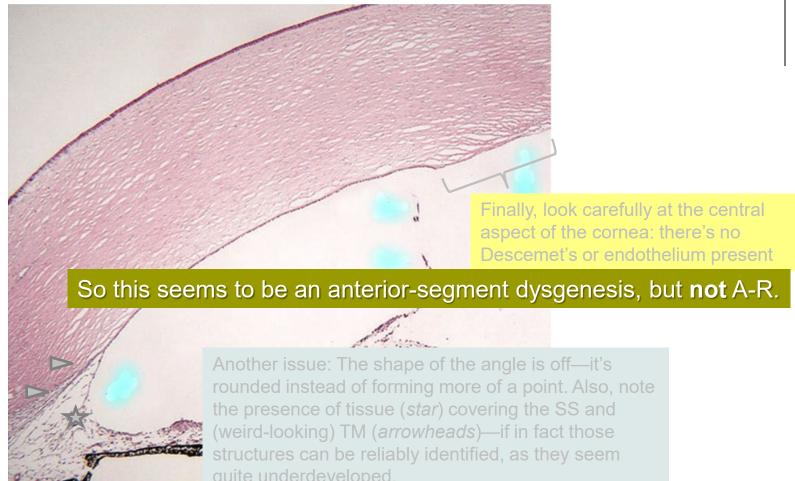










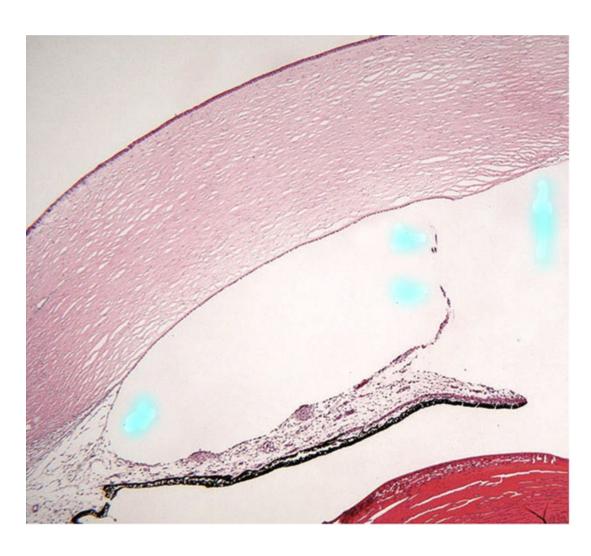




What's the diagnosis?

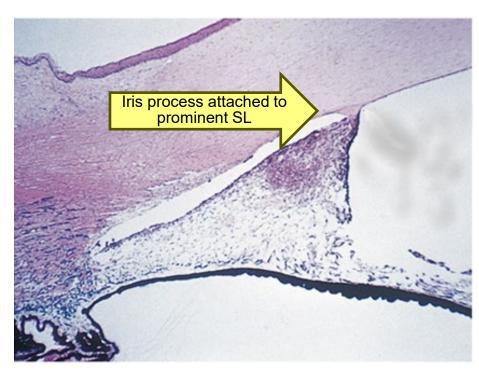


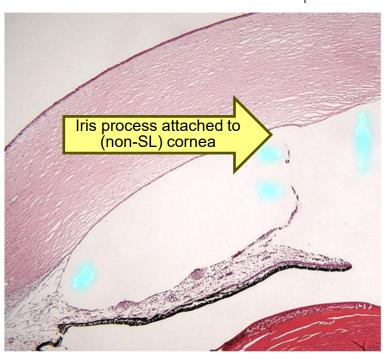




Peter's anomaly. The iris strand attaches to the posterior aspect of the cornea, the central portion of which is missing Descemet's and endothelium. The drainage angle is abnormal, having failed to cleave completely/normally.





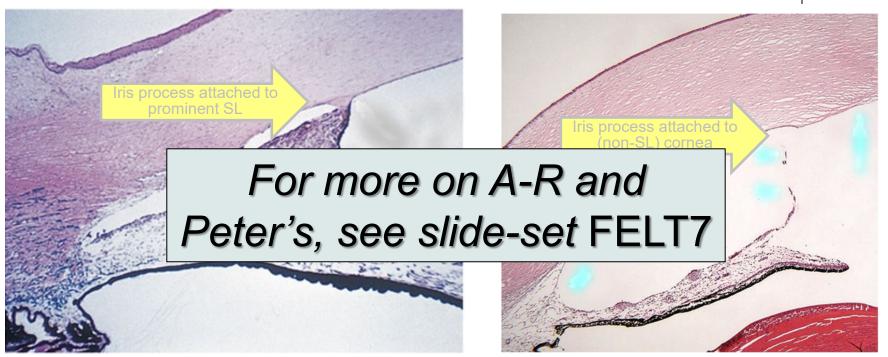


Axenfeld-Rieger

Peter's anomaly

Axenfeld-Rieger and Peter's anomaly side-by-side



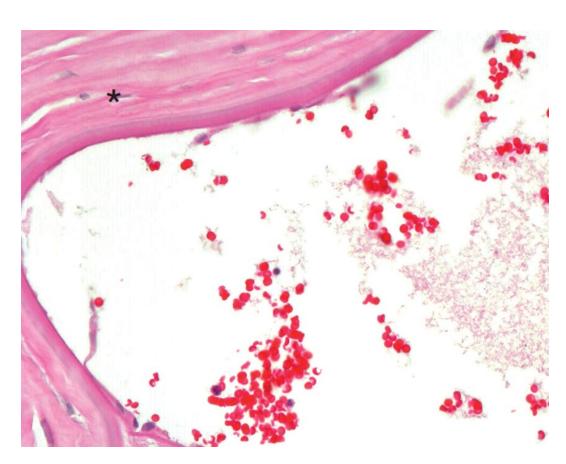


Axenfeld-Rieger

Peter's anomaly

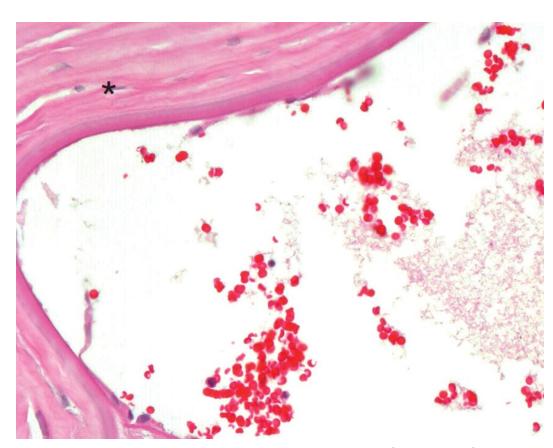
Axenfeld-Rieger and Peter's anomaly side-by-side





As for slides depicting **cell in the AC**: --If the cells are bright red, that's a





Erythrocytes in the anterior chamber (hyphema) posterior to the cornea (asterisk)

As for slides depicting **cell in the AC**: --If the cells are bright red, that's a hyphema



As for slides depicting **cell in the AC**: --If the cells are bright red, that's a hyphema

If the cells are **not** bright red, they are likely specific cell type.



As for slides depicting **cell in the AC**:
--If the cells are bright red, that's a
hyphema

If the cells are **not** bright red, they are likely macrophages .





As for slides depicting **cell in the AC**:
--If the cells are bright red, that's a
hyphema

If the cells are **not** bright red, they are likely macrophages .

--If the macrophages are jet black, you're dealing with a diagnosis





As for slides depicting **cell in the AC**:
--If the cells are bright red, that's a
hyphema

If the cells are **not** bright red, they are likely macrophages .

--If the macrophages are jet black, you're dealing with a melanoma





As for slides depicting **cell in the AC**:
--If the cells are bright red, that's a
hyphema

If the cells are **not** bright red, they are likely macrophages .

--If the macrophages are jet black, you're dealing with a melanoma (that is likely inducing diff diagnosis)



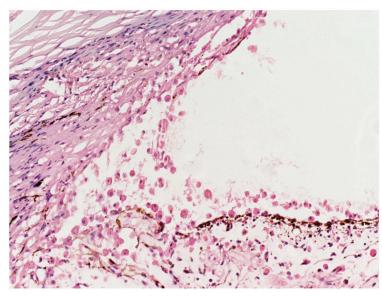


Melanomalytic glaucoma. The trabecular meshwork *(between arrows)* is obstructed by macrophages that have ingested pigment from a necrotic intraocular melanoma.

As for slides depicting **cell in the AC**:
--If the cells are bright red, that's a
hyphema

If the cells are **not** bright red, they are likely macrophages.

--If the macrophages are jet black, you're dealing with a melanoma (that is likely inducing glaucoma)







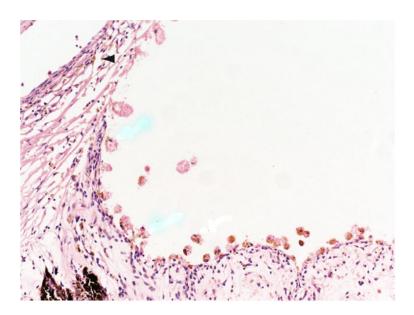
As for slides depicting **cell in the AC**:
--If the cells are bright red, that's a
hyphema

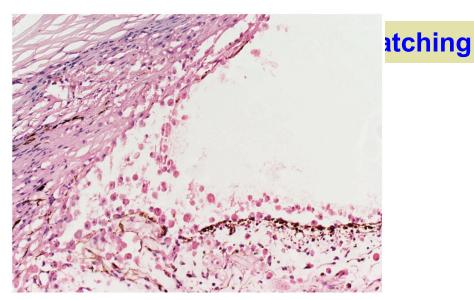
If the cells are **not** bright red, they are likely macrophages.

--If the macrophages are jet black, you're dealing with a melanoma (that is likely inducing glaucoma)

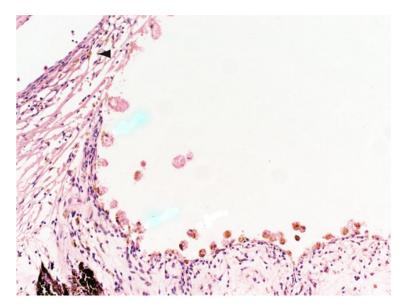
If the macrophages aren't jet black, it's likely one of two entities:

- glaucoma, or - glaucoma





Phacolytic glaucoma showing macrophages filled with degenerated lens cortical material in the angle



Hemolytic glaucoma showing macrophages with erythrocytic debris and hemosiderin in the angle



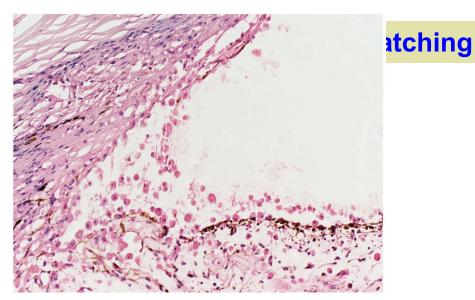
As for slides depicting **cell in the AC**:
--If the cells are bright red, that's a
hyphema

If the cells are **not** bright red, they are likely macrophages.

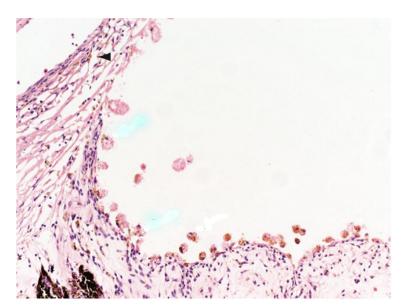
--If the macrophages are jet black, you're dealing with a melanoma (that is likely inducing glaucoma)

If the macrophages aren't jet black, it's likely one of two entities:

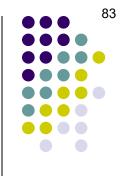
- --Phacolytic glaucoma, or
- --Hemolytic glaucoma



Phacolytic glaucoma showing macrophages filled with degenerated lens cortical material in the angle



Hemolytic glaucoma showing macrophages with erythrocytic debris and hemosiderin in the angle



As for slides depicting **cell in the AC**:
--If the cells are bright red, that's a
hyphema

If the cells are **not** bright red, they are likely macrophages.

--If the macrophages are jet black, you're dealing with a melanoma (that is likely inducing glaucoma)

If the macrophages aren't jet black, it's likely one of two entities:

- --Phacolytic glaucoma, or
- --Hemolytic glaucoma

(It's not apparent to me that these can be reliably distinguished on the basis of appearance, so I suspect clinical context will play a role in doing so.)





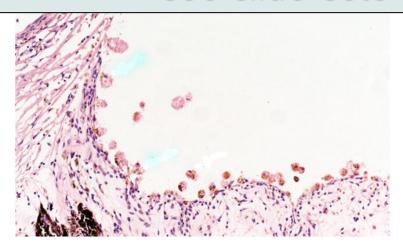
Phacolytic glaucoma showing macrophages filled

As for slides depicting **cell in the AC**:
--If the cells are bright red, that's a
hyphema

If the cells are **not** bright red, they are likely macrophages.

If the magraphages are jet block

For more on phacolytic and hemolytic glaucoma, see slide-sets G13 and G14



Hemolytic glaucoma showing macrophages with erythrocytic debris and hemosiderin in the angle

it's likely one of two entities:

- --Phacolytic glaucoma, or
- --Hemolytic glaucoma

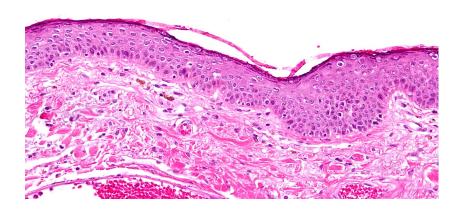
(It's not apparent to me that these can be reliably distinguished on the basis of appearance, so I suspect clinical context will play a role in doing so.)



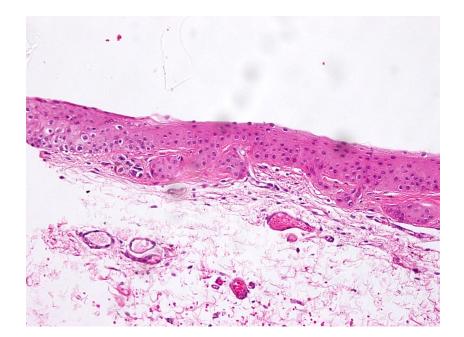
Conj and Lid Skin

These images will include a tissue edge—a boundary between tissue and nothing. Identifying pathology on such images requires that one first determine which tissue (lid skin vs conj) one is dealing with, so we'll start by tackling how to make this distinction.





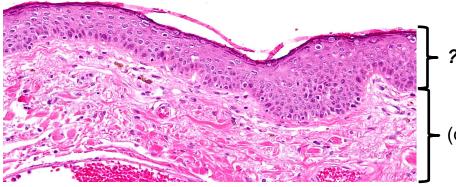
(Photomicrographs courtesy of Dr. Nick Mamalis and his lab)



Here we have conj and lid skin—but which is which?

(Rhetorical question—keep going)

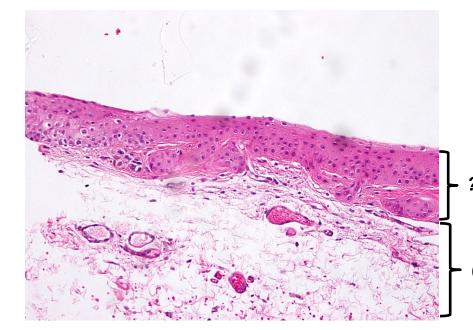




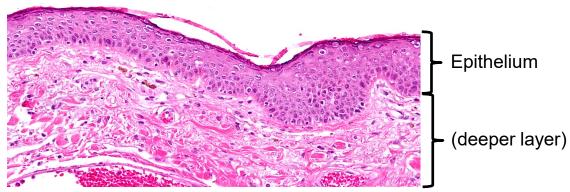
- (deeper layer)

Here we have conj and lid skin—but which is which?

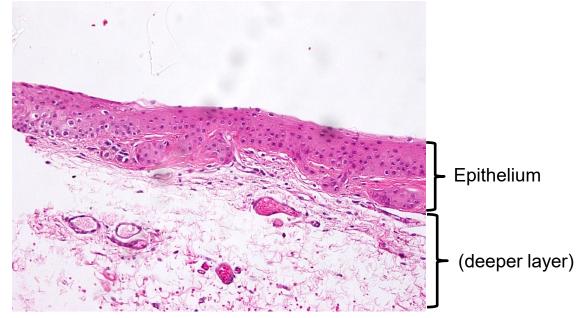
Both consist of two broad layers: An over a deeper layer (to be named shortly).



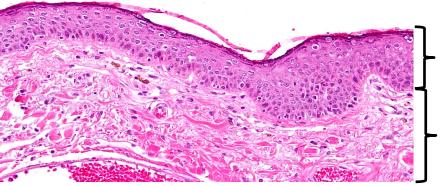




Here we have conj and lid skin—but which is which?
Both consist of two broad layers: An epithelium over a deeper layer (to be named shortly).







Epithelium: cell shape

(deeper layer)

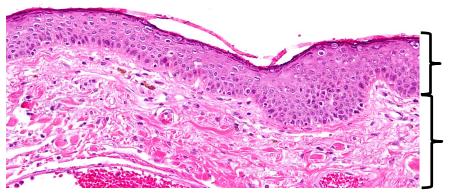
epithelium is cell shape

Epithelium: cell shape

(deeper layer)

Here we have conj and lid skin—but which is which?
Both consist of two broad layers: An epithelium over a deeper layer (to be named shortly). In both, the epithelium is cell shape

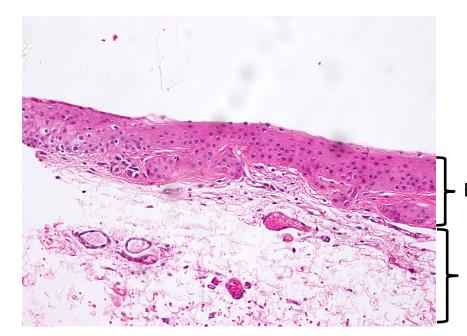




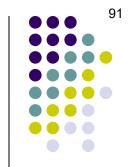
Epithelium: Squamous

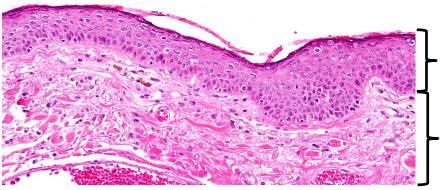
(deeper layer)

Here we have conj and lid skin—but which is which?
Both consist of two broad layers: An epithelium over a deeper layer (to be named shortly). In both, the epithelium is squamous



Epithelium: Squamous





Epithelium: Squamous,

layered-ness

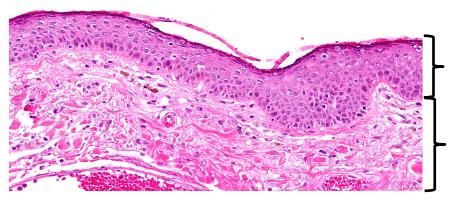
(deeper layer)

Epithelium: Squamous, (deeper layer)

Here we have conj and lid skin—but which is which? Both consist of two broad layers: An epithelium over a deeper layer (to be named shortly). In both, the epithelium is squamous, and

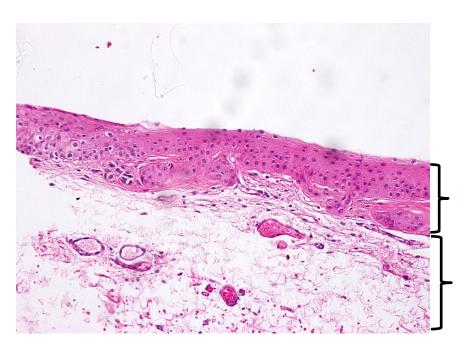
layered-ness





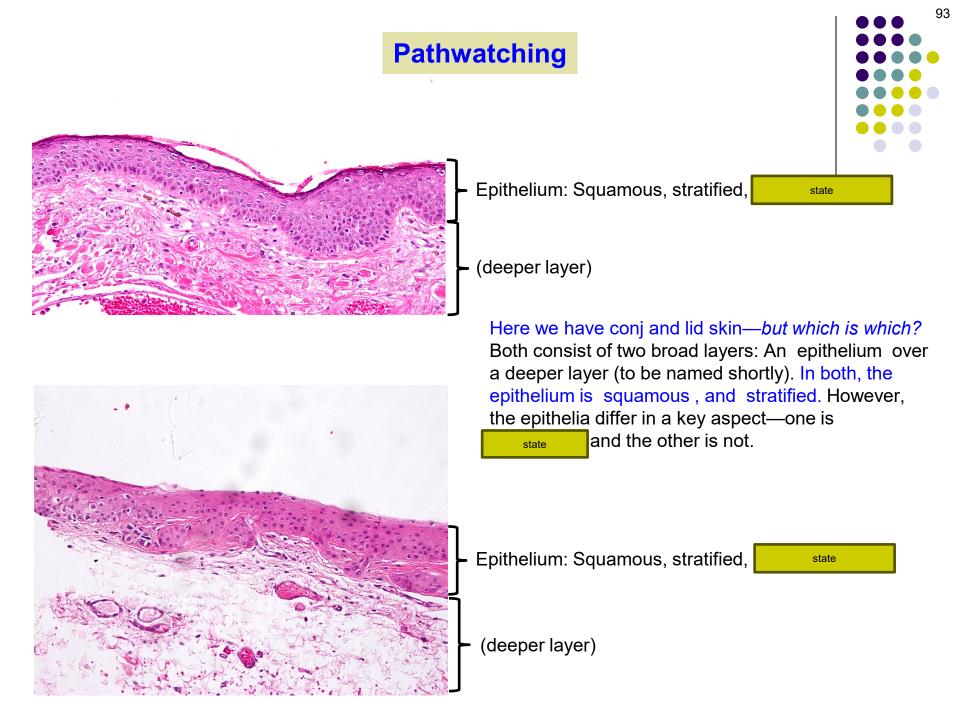
Epithelium: Squamous, stratified

(deeper layer)

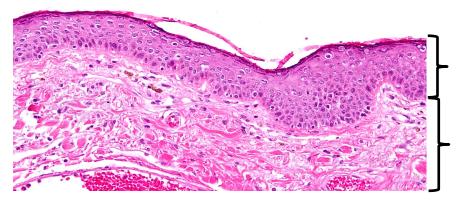


Here we have conj and lid skin—but which is which?
Both consist of two broad layers: An epithelium over a deeper layer (to be named shortly). In both, the epithelium is squamous, and stratified.

Epithelium: Squamous, stratified

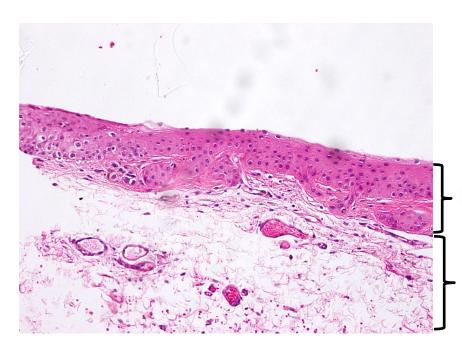






Epithelium: Squamous, stratified, keratinized

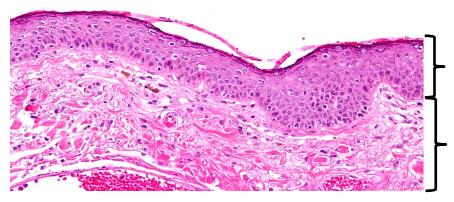
(deeper layer)



Here we have conj and lid skin—but which is which?
Both consist of two broad layers: An epithelium over a deeper layer (to be named shortly). In both, the epithelium is squamous, and stratified. However, the epithelia differ in a key aspect—one is keratinized and the other is not.

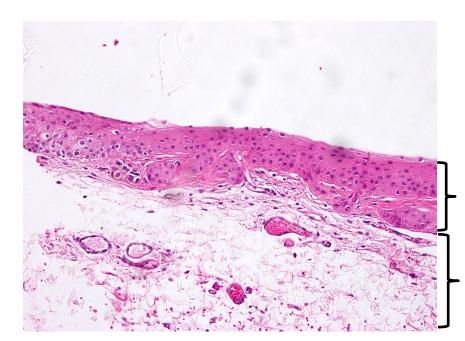
Epithelium: Squamous, stratified, non-keratinized





Epithelium: Squamous, stratified, keratinized

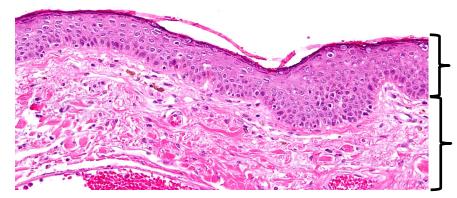
(deeper layer)



Here we have conj and lid skin—but which is which?
Both consist of two broad layers: An epithelium over a deeper layer (to be named shortly). In both, the epithelium is squamous, and stratified. However, the epithelia differ in a key aspect—one is keratinized and the other is not. This is the distinction that allows us to identify them, because one is keratinized, whereas (normal) the other isn't.

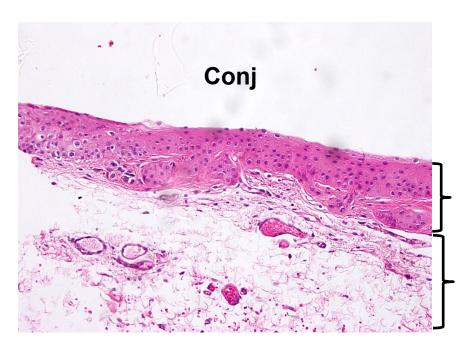
Epithelium: Squamous, stratified, non-keratinized

Lid skin



Epithelium: Squamous, stratified, keratinized

(deeper layer)



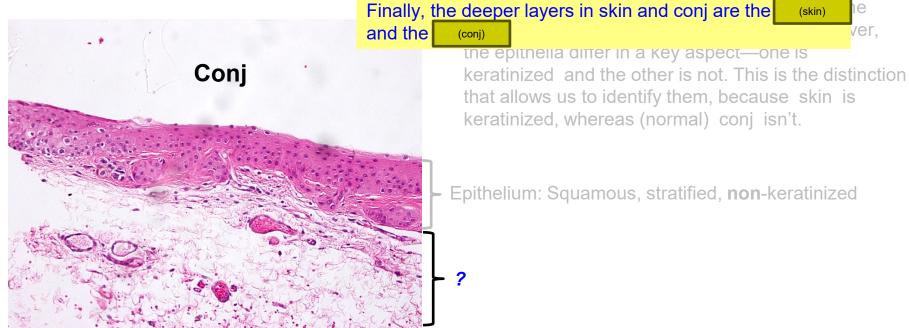
Here we have conj and lid skin—but which is which? Both consist of two broad layers: An epithelium over a deeper layer (to be named shortly). In both, the epithelium is squamous, and stratified. However, the epithelia differ in a key aspect—one is keratinized and the other is not. This is the distinction that allows us to identify them, because skin is keratinized, whereas (normal) conj isn't.

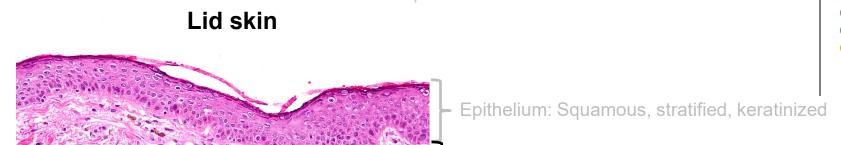
Epithelium: Squamous, stratified, non-keratinized



Here we have conj and lid skin—but which is which?
Both consist of two broad layers: An epithelium over

97





Dermis

Here we have conj and lid skin—but which is which?
Both consist of two broad layers: An epithelium over

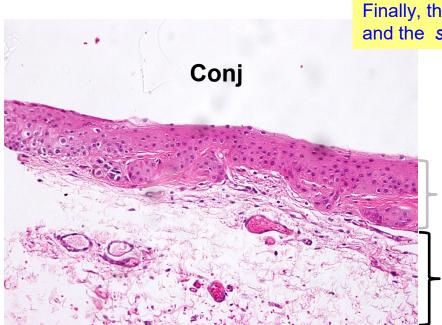
98

Finally, the deeper layers in skin and conj are the *dermis* and the *stroma*

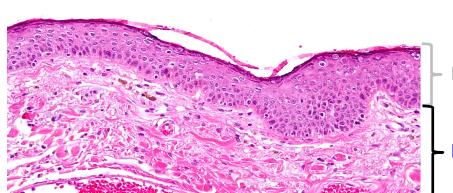
tne epitnella diπer in a key aspect—one is keratinized and the other is not. This is the distinction that allows us to identify them, because skin is keratinized, whereas (normal) conj isn't.

Epithelium: Squamous, stratified, non-keratinized

Stroma



Lid skin



Epithelium: Squamous, stratified, keratinized

Dermis

Here we have conj and lid skin—but which is which? Both consist of two broad layers: An epithelium over

99

Finally, the deeper layers in skin and conj are the *dermis* and the *stroma* (aka Conj

) respectively two words tne epitnella differ in a key aspect—one is keratinized and the other is not. This is the distinction that allows us to identify them, because skin is keratinized, whereas (normal) conj isn't.

Epithelium: Squamous, stratified, non-keratinized

Stroma or

two words



Dermis

Here we have conj and lid skin—but which is which?
Both consist of two broad layers: An epithelium over

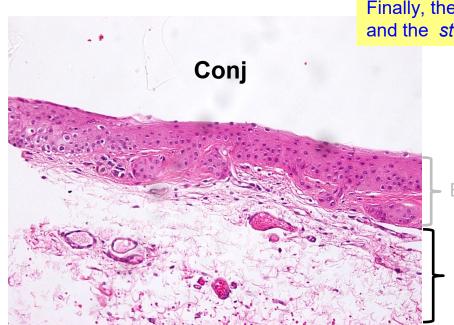
100

Finally, the deeper layers in skin and conj are the *dermis* and the *stroma* (aka *substantia propia*) respectively

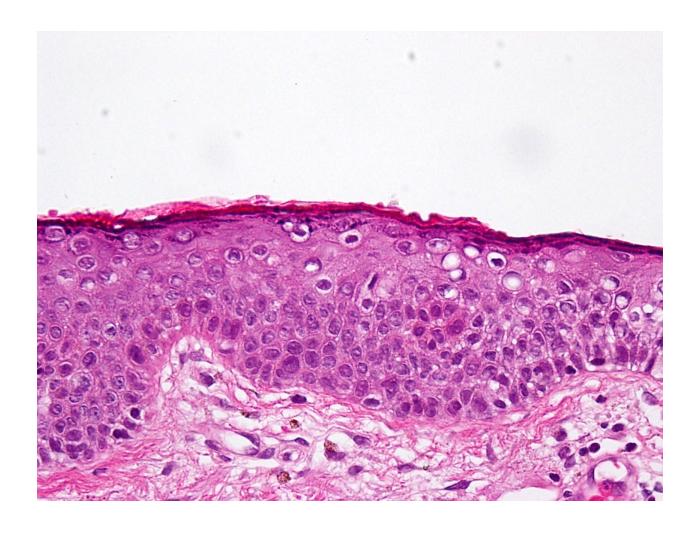
the epithelia differ in a key aspect—one is keratinized and the other is not. This is the distinction that allows us to identify them, because skin is keratinized, whereas (normal) conj isn't.

Epithelium: Squamous, stratified, non-keratinized

Stroma or substantia propia

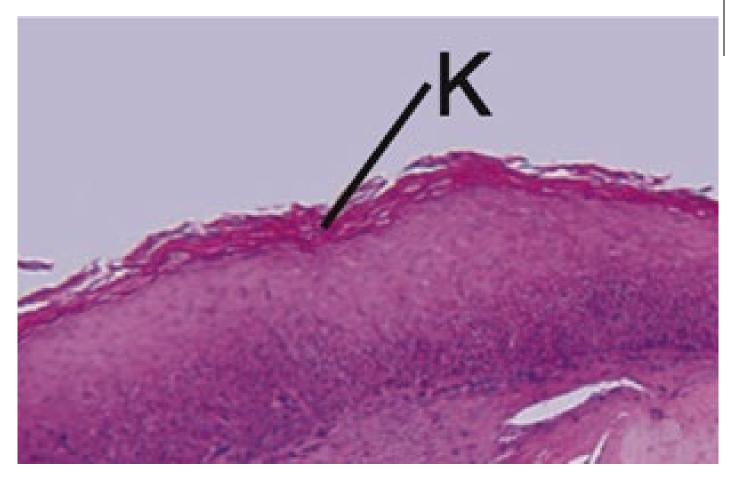






Note 1: Lid keratinization can be subtler than depicted previously





Note 2: Keratinization occurs in some conj pathologies (OSSN* in this case)

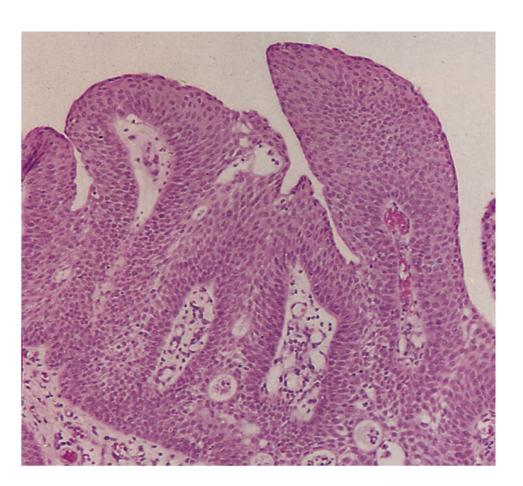
*Which stands for





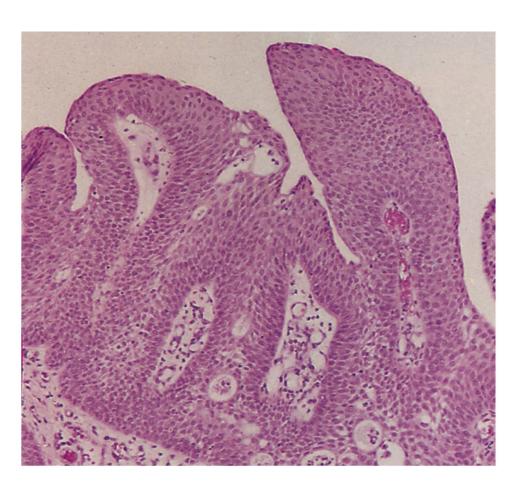
Note 2: Keratinization occurs in some conj pathologies (OSSN* in this case)





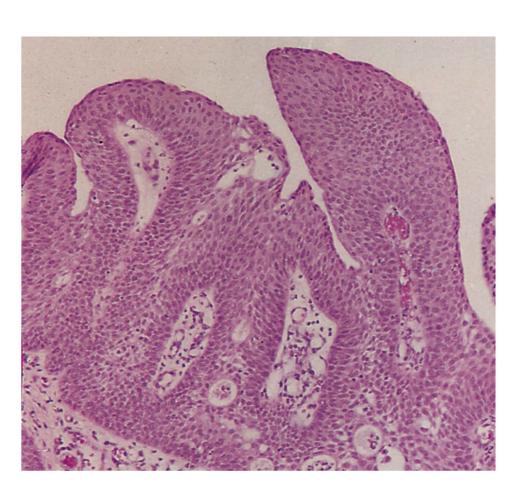
First things first: Skin, or conj?





First things first: Skin, or conj?
Lack of keratinization = conj

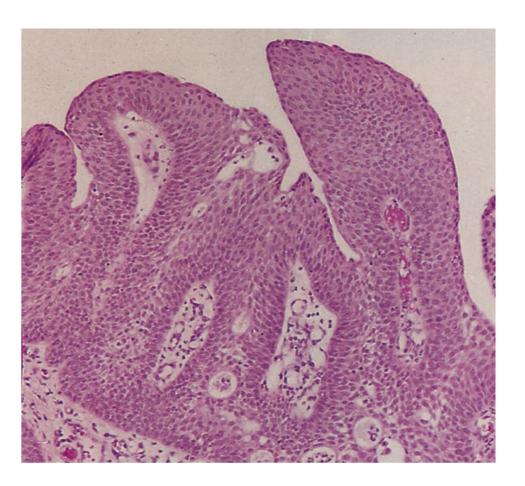




First things first: Skin, or conj?
Lack of keratinization = conj

Clearly this is not normal conj—there is obvious exuberant growth, indicating it's a ______.

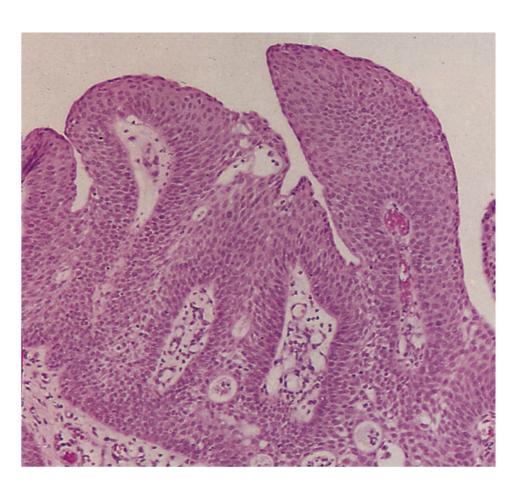




First things first: Skin, or conj?
Lack of keratinization = conj

Clearly this is not normal conj—there is obvious exuberant growth, indicating it's a neoplasia*.





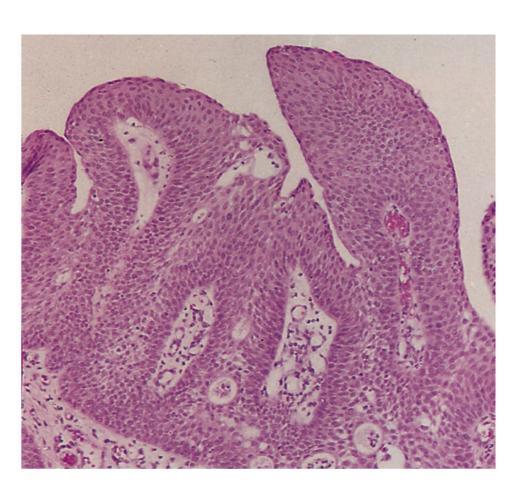
First things first: Skin, or conj?
Lack of keratinization = conj

Clearly this is not normal conj—there is obvious exuberant growth, indicating it's a neoplasia*.

But how to begin identifying it?

Begin by recognizing there are # broad categories of conj neoplasias:



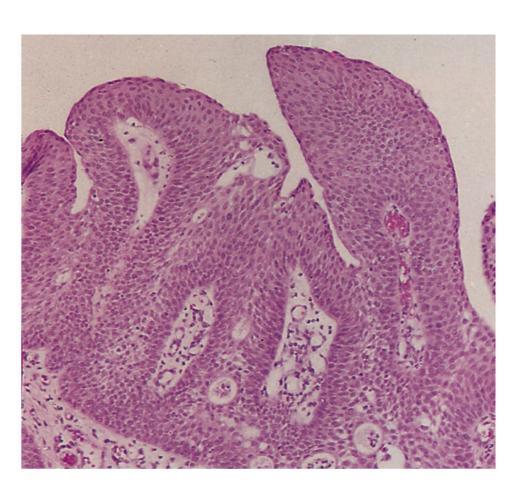


First things first: Skin, or conj?
Lack of keratinization = conj

Clearly this is not normal conj—there is obvious exuberant growth, indicating it's a neoplasia*. But how to begin identifying it?
Begin by recognizing there are three broad categories of conj neoplasias:

- --?
- --?
- --?





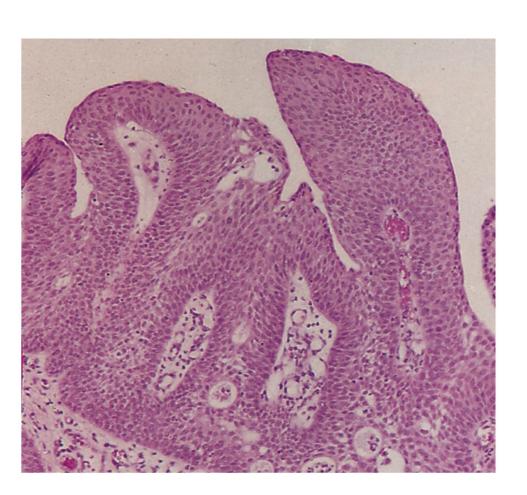
First things first: Skin, or conj?
Lack of keratinization = conj

Clearly this is not normal conj—there is obvious exuberant growth, indicating it's a neoplasia*. But how to begin identifying it?

Begin by recognizing there are three broad categories of conj neoplasias:

- --Lymphatic
- --Melanocytic
- --Epithelial





First things first: Skin, or conj?
Lack of keratinization = conj

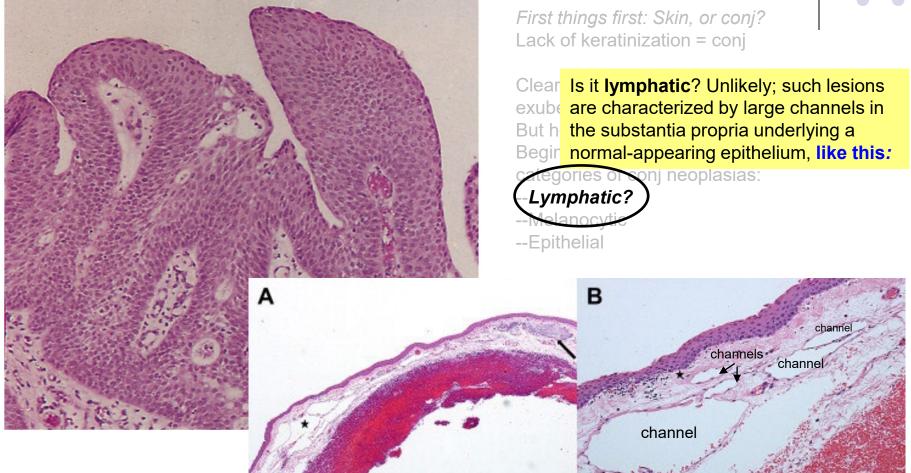
Clear Is it Iymphatic? Unlikely; such lesions exuberare characterized by large channels in the substantia propria underlying a Begin normal-appearing epithelium

categories of sonj neoplasias:

Lymphatic?

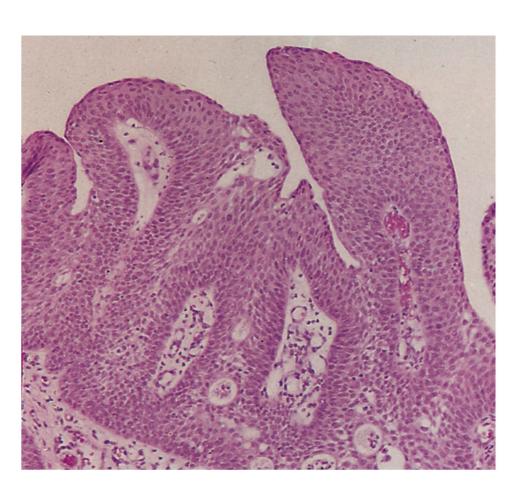
- --Melanocyti
- --Epithelial





Lymphangiectasia. *A*, low power; *B*, higher power





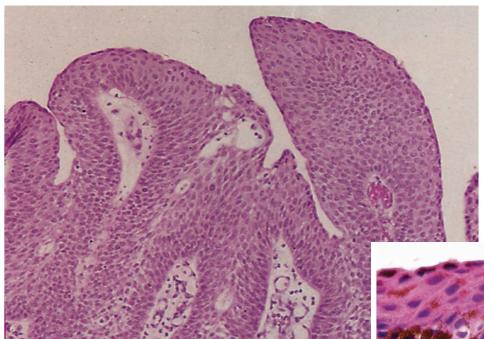
First things first: Skin, or conj?
Lack of keratinization = conj

Clear Is it **melanocytic**? Also unlikely, as such exube lesions typically contain an attention-But he grabbing amount of melanin Begin

categories of conj neoplasias:



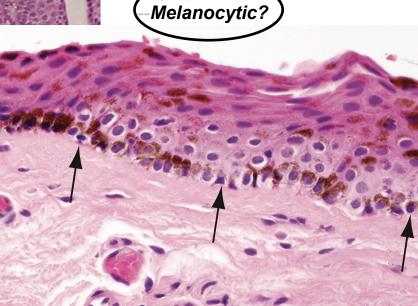




First things first: Skin, or conj?
Lack of keratinization = conj

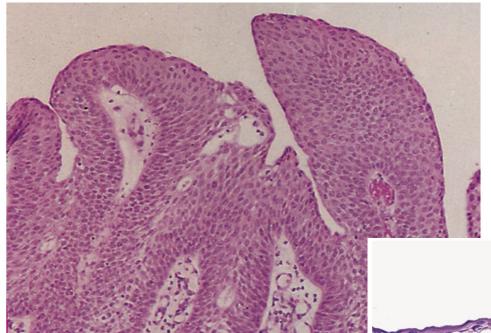
Clear Is it **melanocytic**? Also unlikely, as such exube lesions typically contain an attention-grabbing amount of melanin like **this**Begin

categories of conj neoplasias:



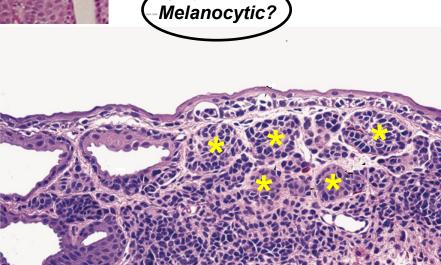
Complexion-associated (aka racial, aka benign-acquired) melanosis





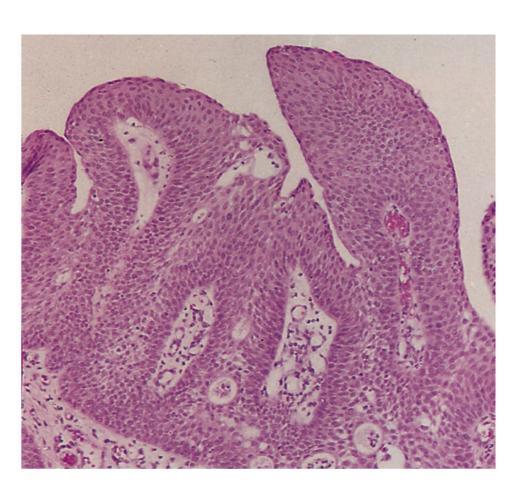
First things first: Skin, or conj?
Lack of keratinization = conj

Clear Is it **melanocytic**? Also unlikely, as such lesions typically contain an attention-grabbing amount of melanin like **this**, and/or lots of melanocytes like **this** categories of conj neoplasias:



Conj nevus with beaucoup melanocytes in nests (asterisks)





First things first: Skin, or conj?
Lack of keratinization = conj

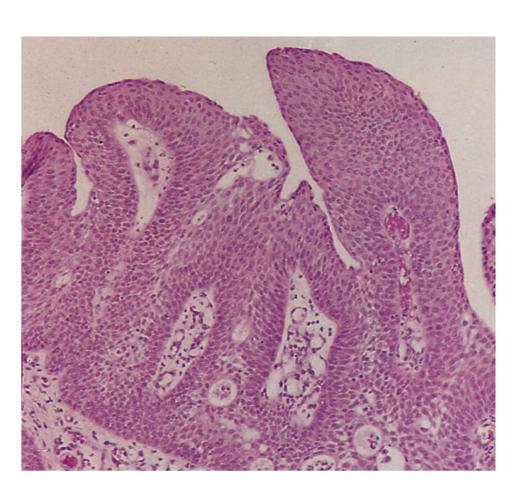
Clearly this is not normal conj—there is obvious exuberant growth, indicating it's a neoplasia. But how to begin identifying it?

Begin by recognizing there are three broad categories of conj neoplasias:

- --Lymphatic
- --Melanocytic
- --Epithelial

So by process of elimination, it's epithelial.





First things first: Skin, or conj?
Lack of keratinization = conj

Clearly this is not normal conj—there is obvious exuberant growth, indicating it's a neoplasia. But how to begin identifying it?

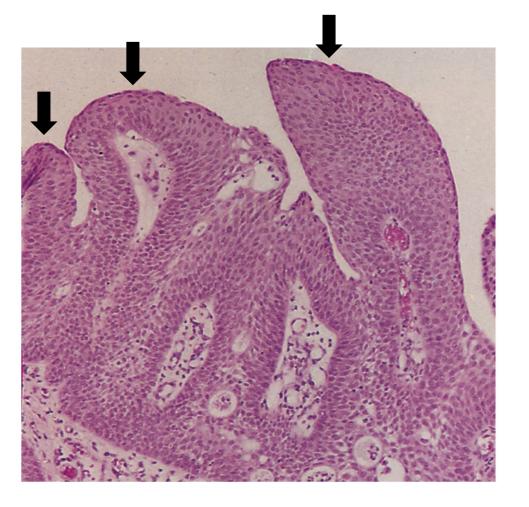
Begin by recognizing there are three broad categories of conj neoplasias:

- --Lymphatic
- --Melanocytic
- --Epithelial

So by process of elimination, it's **epithelial**.

But what is it? In this case, there's a classic field mark:





First things first: Skin, or conj?
Lack of keratinization = conj

Clearly this is not normal conj—there is obvious exuberant growth, indicating it's a neoplasia. But how to begin identifying it?

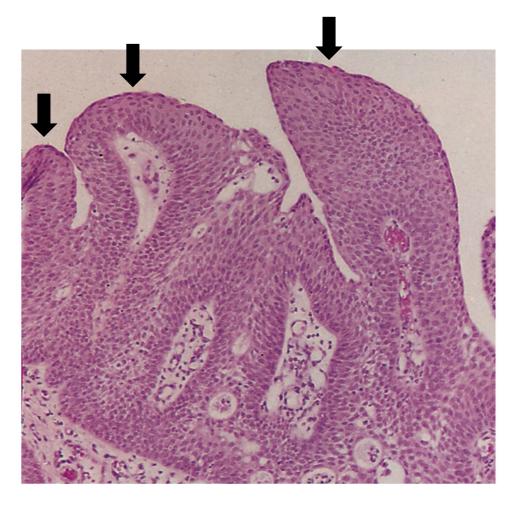
Begin by recognizing there are three broad categories of conj neoplasias:

- --Lymphatic
- --Melanocytic
- --Epithelial

So by process of elimination, it's **epithelial**.

But what is it? In this case, there's a classic field mark: **These projections**, classically described as _____.





First things first: Skin, or conj?
Lack of keratinization = conj

Clearly this is not normal conj—there is obvious exuberant growth, indicating it's a neoplasia. But how to begin identifying it?

Begin by recognizing there are three broad categories of conj neoplasias:

- --Lymphatic
- --Melanocytic
- --Epithelial

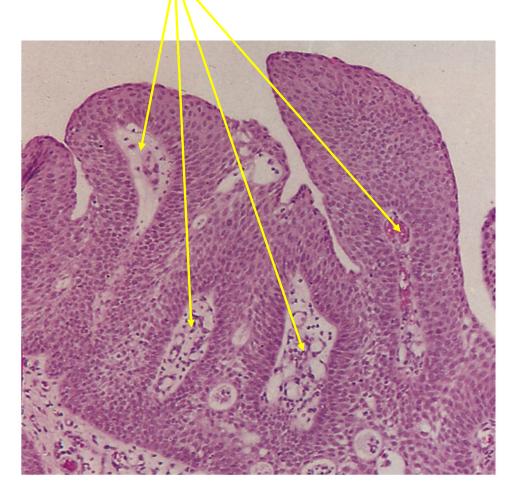
So by process of elimination, it's **epithelial**.

But what is it? In this case, there's a classic field mark: **These projections**, classically described as 'fronds'.

Note that the 'cores' of the fronds are structures

Pathwatching





First things first: Skin, or conj?
Lack of keratinization = conj

Clearly this is not normal conj—there is obvious exuberant growth, indicating it's a neoplasia. But how to begin identifying it?

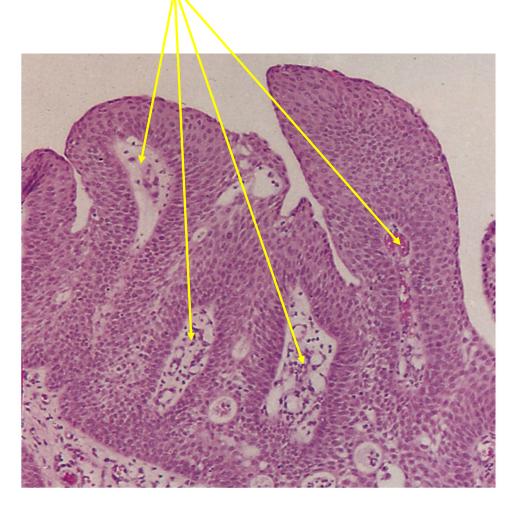
Begin by recognizing there are three broad categories of conj neoplasias:

- --Lymphatic
- --Melanocytic
- --Epithelial

So by process of elimination, it's **epithelial**.

But what is it? In this case, there's a classic field mark: **These projections**, classically described as 'fronds'.





First things first: Skin, or conj?
Lack of keratinization = conj

Clearly this is not normal conj—there is obvious exuberant growth, indicating it's a neoplasia. But how to begin identifying it?

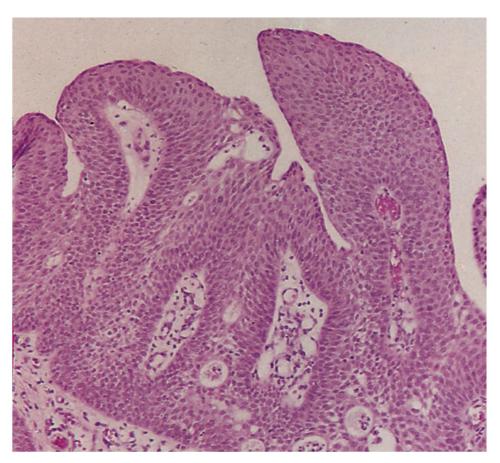
Begin by recognizing there are three broad categories of conj neoplasias:

- --Lymphatic
- --Melanocytic
- --Epithelial

So by process of elimination, it's **epithelial**.

But what is it? In this case, there's a classic field mark: **These projections**, classically described as 'fronds'.





What's the diagnosis?

First things first: Skin, or conj?
Lack of keratinization = conj

Clearly this is not normal conj—there is obvious exuberant growth, indicating it's a neoplasia. But how to begin identifying it?

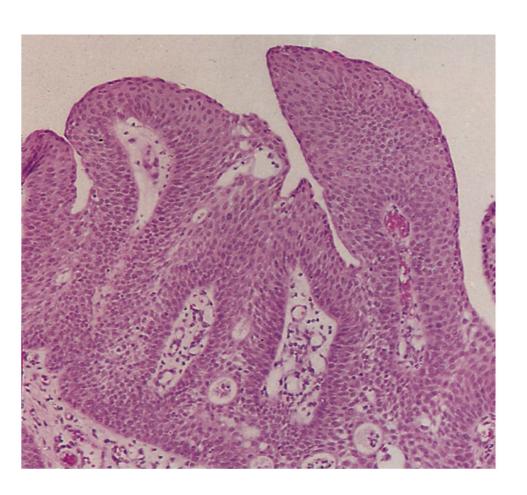
Begin by recognizing there are three broad categories of conj neoplasias:

- --Lymphatic
- --Melanocytic
- --Epithelial

So by process of elimination, it's **epithelial**.

But what is it? In this case, there's a classic field mark: **These projections**, classically described as 'fronds'. When you see a conj lesion with a 'frond' appearance, one term should come to mind:





First things first: Skin, or conj? Lack of keratinization = conj

Clearly this is not normal conj—there is obvious exuberant growth, indicating it's a neoplasia. But how to begin identifying it? Begin by recognizing there are three broad categories of conj neoplasias:

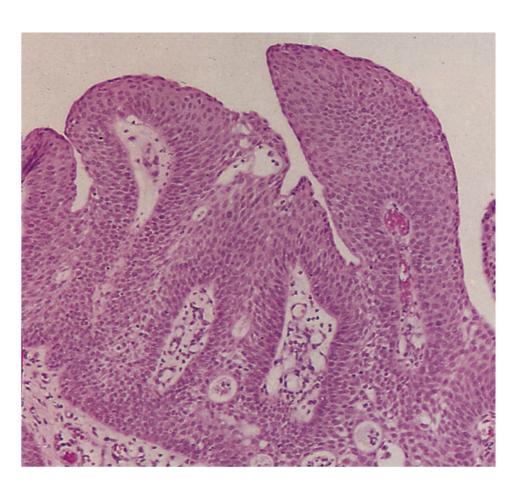
- --Lymphatic
- --Melanocytic
- --Epithelial

So by process of elimination, it's epithelial.

But what is it? In this case, there's a classic field mark: **These projections**, classically described as 'fronds'. When you see a conj lesion with a 'frond' appearance, one term should come to mind:

Pedunculated papilloma of the conj occur more often in adults vides





First things first: Skin, or conj?
Lack of keratinization = conj

Clearly this is not normal conj—there is obvious exuberant growth, indicating it's a neoplasia. But how to begin identifying it?

Begin by recognizing there are three broad categories of conj neoplasias:

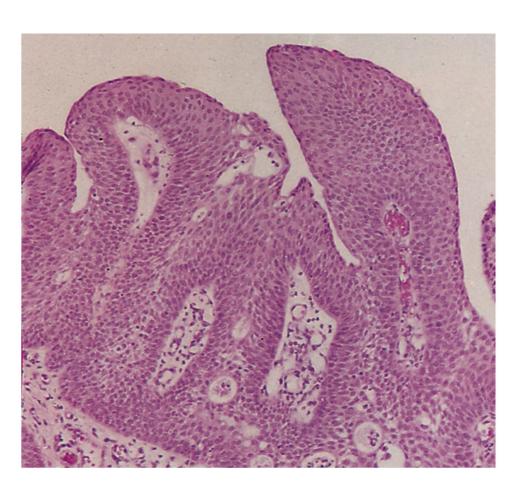
- --Lymphatic
- --Melanocytic
- --Epithelial

So by process of elimination, it's **epithelial**.

But what is it? In this case, there's a classic field mark: **These projections**, classically described as 'fronds'. When you see a conj lesion with a 'frond' appearance, one term should come to mind:

Pedunculated papilloma of the conj occur more often in kids .





First things first: Skin, or conj?
Lack of keratinization = conj

Clearly this is not normal conj—there is obvious exuberant growth, indicating it's a neoplasia. But how to begin identifying it?

Begin by recognizing there are three broad categories of conj neoplasias:

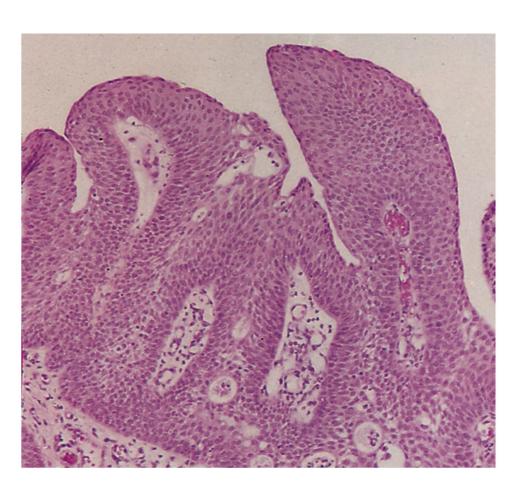
- --Lymphatic
- --Melanocytic
- --Epithelial

So by process of elimination, it's epithelial.

But what is it? In this case, there's a classic field mark: **These projections**, classically described as 'fronds'. When you see a conj lesion with a 'frond' appearance, one term should come to mind:

Pedunculated papilloma of the conj occur more often in kids . They are associated with certain subtypes of labb. infection.





First things first: Skin, or conj?
Lack of keratinization = conj

Clearly this is not normal conj—there is obvious exuberant growth, indicating it's a neoplasia. But how to begin identifying it?

Begin by recognizing there are three broad categories of conj neoplasias:

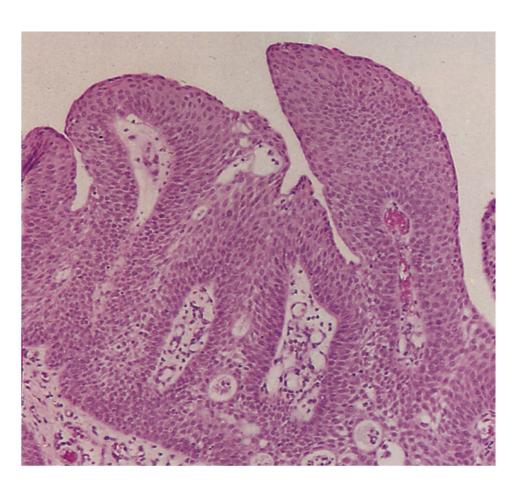
- --Lymphatic
- --Melanocytic
- --Epithelial

So by process of elimination, it's epithelial.

But what is it? In this case, there's a classic field mark: **These projections**, classically described as 'fronds'. When you see a conj lesion with a 'frond' appearance, one term should come to mind:

Pedunculated papilloma of the conj occur more often in kids . They are associated with certain subtypes of HPV infection.





First things first: Skin, or conj?
Lack of keratinization = conj

Clearly this is not normal conj—there is obvious exuberant growth, indicating it's a neoplasia. But how to begin identifying it?

Begin by recognizing there are three broad categories of conj neoplasias:

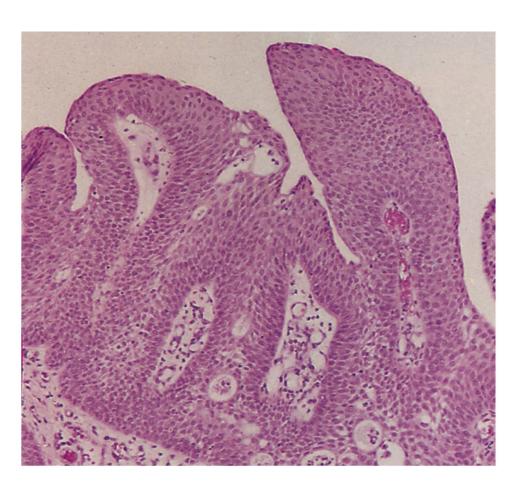
- --Lymphatic
- --Melanocytic
- --Epithelial

So by process of elimination, it's epithelial.

But what is it? In this case, there's a classic field mark: **These projections**, classically described as 'fronds'. When you see a conj lesion with a 'frond' appearance, one term should come to mind:

Pedunculated papilloma of the conj occur more often in kids . They are associated with certain subtypes of HPV infection. They have significant vs no malignant potential.





First things first: Skin, or conj?
Lack of keratinization = conj

Clearly this is not normal conj—there is obvious exuberant growth, indicating it's a neoplasia. But how to begin identifying it?

Begin by recognizing there are three broad categories of conj neoplasias:

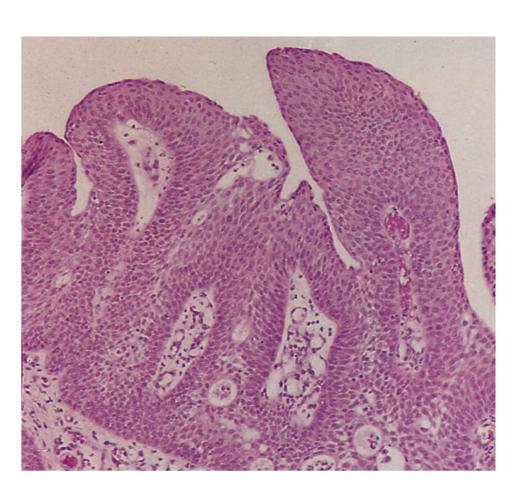
- --Lymphatic
- --Melanocytic
- --Epithelial

So by process of elimination, it's epithelial.

But what is it? In this case, there's a classic field mark: **These projections**, classically described as 'fronds'. When you see a conj lesion with a 'frond' appearance, one term should come to mind:

Pedunculated papilloma of the conj occur more often in kids . They are associated with certain subtypes of HPV infection. They have negligible malignant potential.





First things first: Skin, or conj?
Lack of keratinization = conj

Clearly this is not normal conj—there is obvious exuberant growth, indicating it's a neoplasia. But how to begin identifying it?

Begin by recognizing there are three broad categories of conj neoplasias:

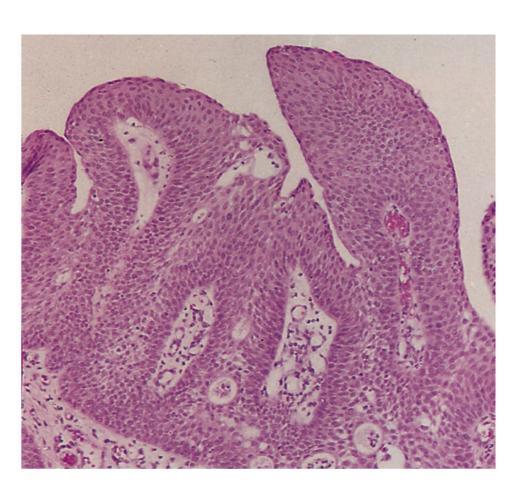
- --Lymphatic
- --Melanocytic
- --Epithelial

So by process of elimination, it's **epithelial**.

But what is it? In this case, there's a classic field mark: **These projections**, classically described as 'fronds'. When you see a conj lesion with a 'frond' appearance, one term should come to mind:

Pedunculated papilloma of the conj occur more often in kids. They are associated with certain subtypes of HPV infection. They have negligible malignant potential. In contrast, papillomas are more common in adults.





First things first: Skin, or conj?
Lack of keratinization = conj

Clearly this is not normal conj—there is obvious exuberant growth, indicating it's a neoplasia. But how to begin identifying it?

Begin by recognizing there are three broad categories of conj neoplasias:

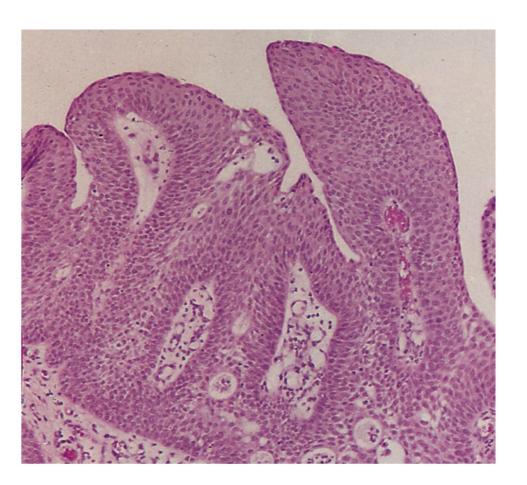
- --Lymphatic
- --Melanocytic
- --Epithelial

So by process of elimination, it's **epithelial**.

But what is it? In this case, there's a classic field mark: **These projections**, classically described as 'fronds'. When you see a conj lesion with a 'frond' appearance, one term should come to mind:

Pedunculated papilloma of the conj occur more often in kids. They are associated with certain subtypes of HPV infection. They have negligible malignant potential. In contrast, sessile papillomas are more common in adults.





First things first: Skin, or conj?
Lack of keratinization = conj

Clearly this is not normal conj—there is obvious exuberant growth, indicating it's a neoplasia. But how to begin identifying it?

Begin by recognizing there are three broad categories of conj neoplasias:

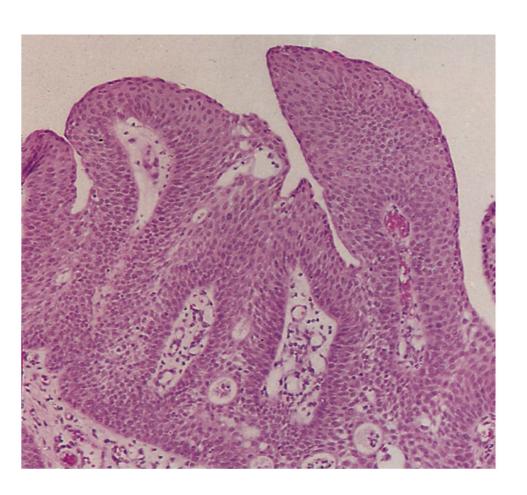
- --Lymphatic
- --Melanocytic
- --Epithelial

So by process of elimination, it's **epithelial**.

But what is it? In this case, there's a classic field mark: **These projections**, classically described as 'fronds'. When you see a conj lesion with a 'frond' appearance, one term should come to mind:

Pedunculated papilloma of the conj occur more often in kids. They are associated with certain subtypes of HPV infection. They have negligible malignant potential. In contrast, sessile papillomas are more common in adults. They also are associated with certain (different) HPV subtypes. Their malignant potential is significant vs negligible.





First things first: Skin, or conj?
Lack of keratinization = conj

Clearly this is not normal conj—there is obvious exuberant growth, indicating it's a neoplasia. But how to begin identifying it?

Begin by recognizing there are three broad categories of conj neoplasias:

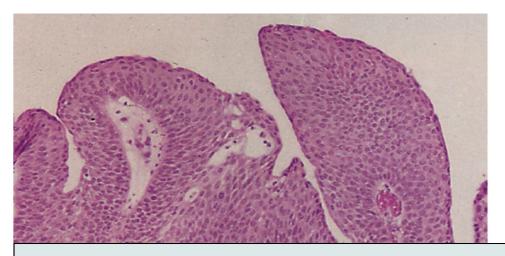
- --Lymphatic
- --Melanocytic
- --Epithelial

So by process of elimination, it's **epithelial**.

But what is it? In this case, there's a classic field mark: **These projections**, classically described as 'fronds'. When you see a conj lesion with a 'frond' appearance, one term should come to mind:

Pedunculated papilloma of the conj occur more often in kids. They are associated with certain subtypes of HPV infection. They have negligible malignant potential. In contrast, sessile papillomas are more common in adults. They also are associated with certain (different) HPV subtypes. Their malignant potential is significant.





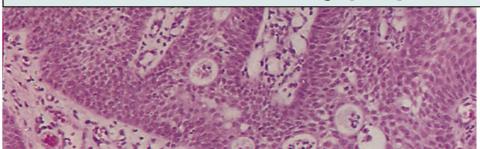
First things first: Skin, or conj? Lack of keratinization = conj

Clearly this is not normal conj—there is obvious exuberant growth, indicating it's a neoplasia. But how to begin identifying it?

Begin by recognizing there are three broad categories of conj neoplasias:

- --Lymphatic
- --Melanocytic

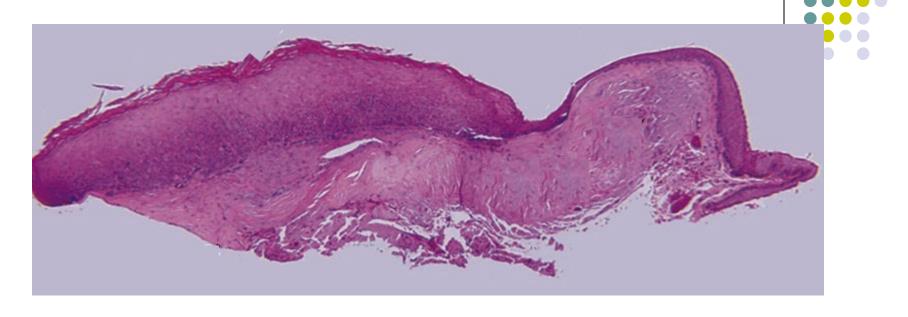
For more on conj papillomas, see slide-set K25



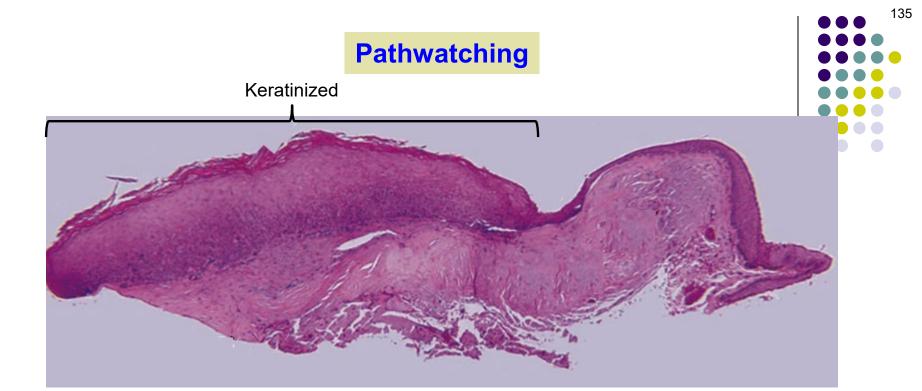
But what is it? In this case, there's a classic field mark: **These projections**, classically described as 'fronds'. When you see a conj lesion with a 'frond' appearance, one term should come to mind:

Pedunculated papilloma of the conj occur more often in kids. They are associated with certain subtypes of HPV infection. They have negligible malignant potential. In contrast, *sessile* papillomas are more common in adults. They also are associated with certain (different) HPV subtypes. Their malignant potential is significant.

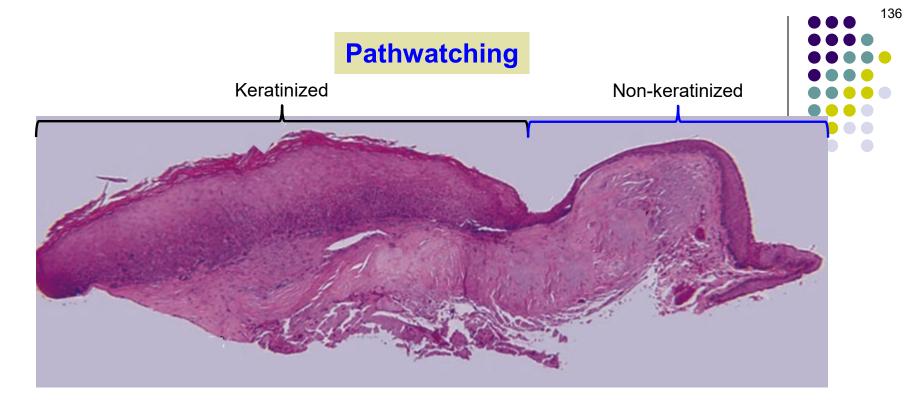
134



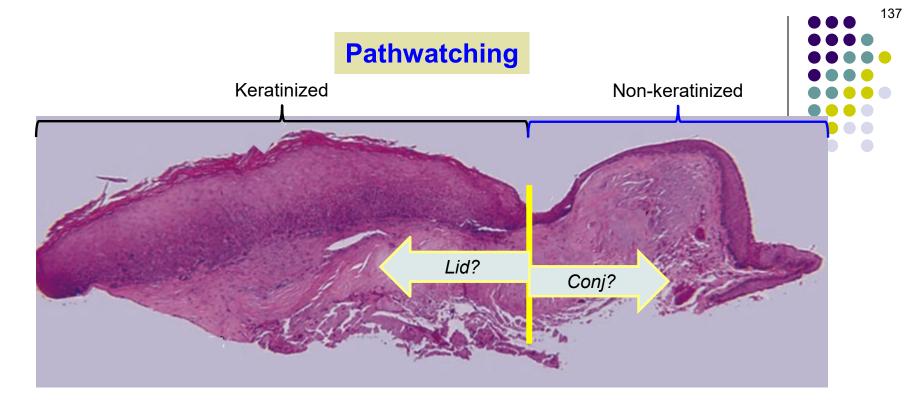
Again, first things first: Skin, or conj?



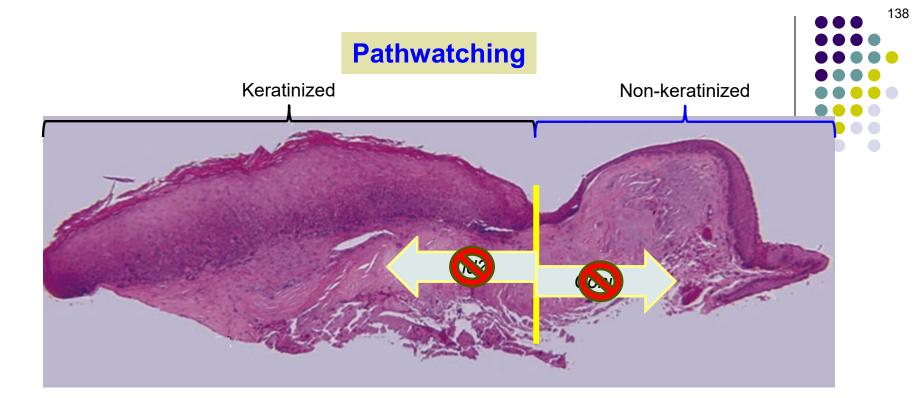
Again, first things first: Skin, or conj?
At first blush this seems an impossible question to answer, because part of it is keratinized (and therefore indicates this is skin)



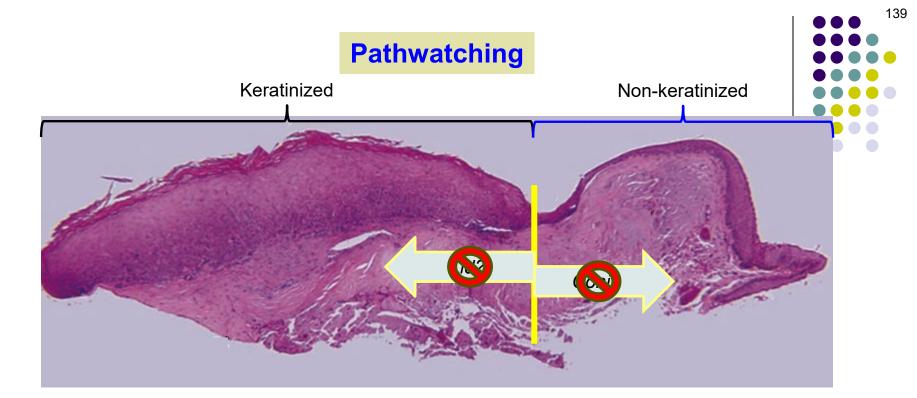
Again, first things first: Skin, or conj?
At first blush this seems an impossible question to answer, because part of it is keratinized (and therefore indicates this is skin) and part of it isn't (indicating conj).



Again, first things first: Skin, or conj?
At first blush this seems an impossible question to answer, because part of it is keratinized (and therefore indicates this is skin) and part of it isn't (indicating conj). Could this be a junction between lid skin and conj?

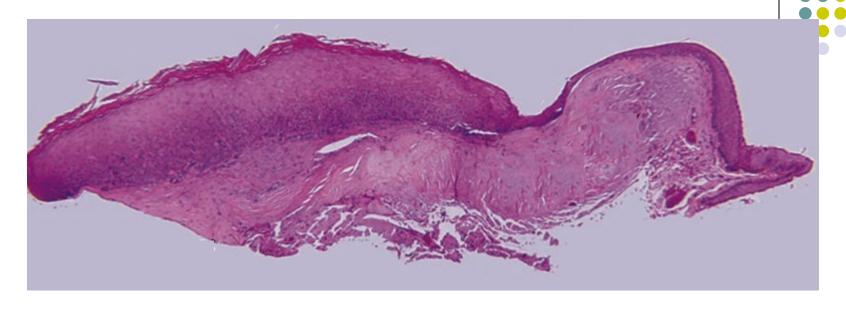


At first blush this seems an impossible question to answer, because part of it is keratinized (and therefore indicates this is skin) and part of it isn't (indicating conj). Could this be a junction between lid skin and conj? No, because no such junction exists (without intervening landmarks of the two words).



Again, first things first: Skin, or conj?

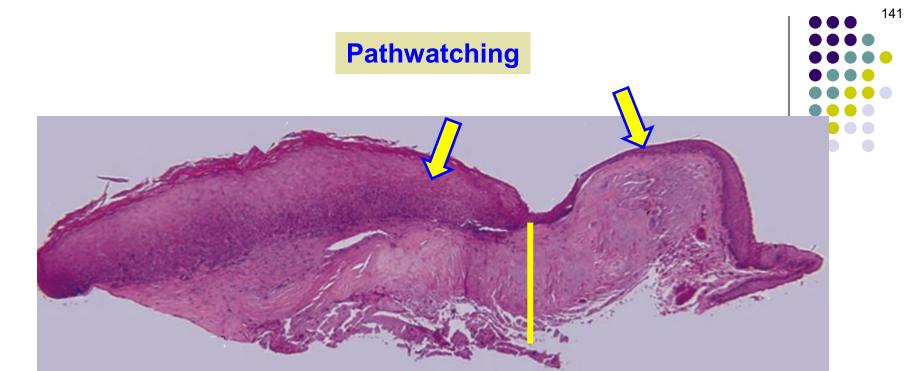
At first blush this seems an impossible question to answer, because part of it is keratinized (and therefore indicates this is skin) and part of it isn't (indicating conj). Could this be a junction between lid skin and conj? No, because no such junction exists (without intervening landmarks of the lid margin).



Again, first things first: Skin, or conj?

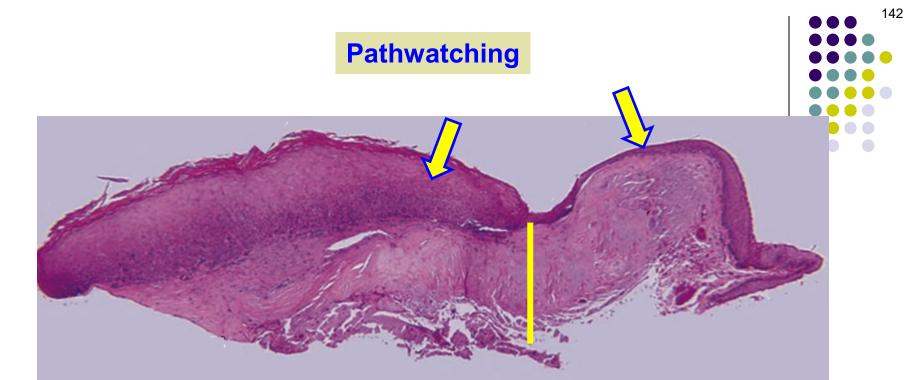
At first blush this seems an impossible question to answer, because part of it is keratinized (and therefore indicates this is skin) and part of it isn't (indicating conj). Could this be a junction between lid skin and conj? No, because no such junction exists (without intervening landmarks of the lid margin). So this is either skin that lost keratinization or conj that acquired it. But which is it?

140



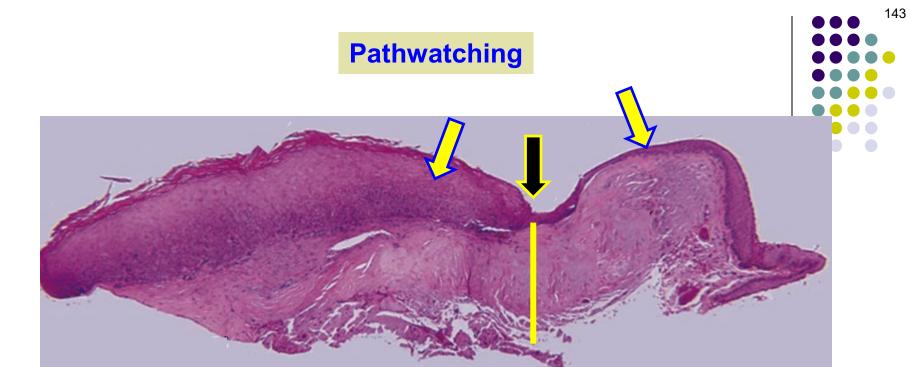
At first blush this seems an impossible question to answer, because part of it is keratinized (and therefore indicates this is skin) and part of it isn't (indicating conj). Could this be a junction between lid skin and conj? No, because no such junction exists (without intervening landmarks of the lid margin). So this is either skin that lost keratinization or conj that acquired it. But which is it?

The answer is clear once we compare the epithelia on the two sides of the change in keratinization status.



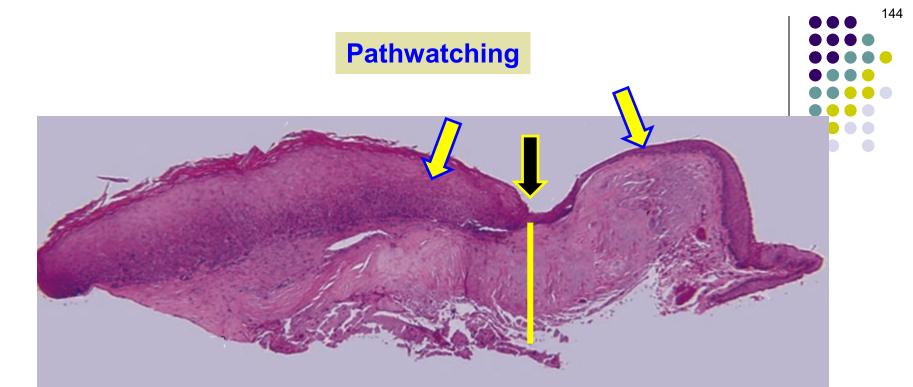
At first blush this seems an impossible question to answer, because part of it is keratinized (and therefore indicates this is skin) and part of it isn't (indicating conj). Could this be a junction between lid skin and conj? No, because no such junction exists (without intervening landmarks of the lid margin). So this is either skin that lost keratinization or conj that acquired it. But which is it?

The answer is clear once we compare the epithelia on the two sides of the change in keratinization status. The epithelium on the keratinized side is vastly thicker than that on the nonkeratinized side, strongly suggesting it (the keratinized epithelium) is neoplastic.



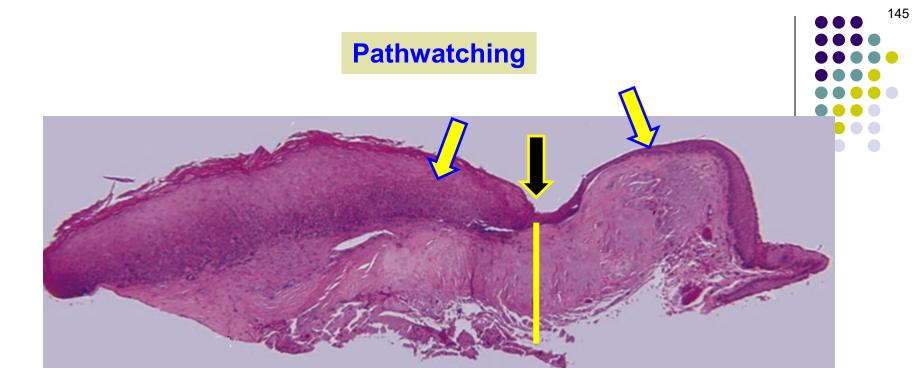
At first blush this seems an impossible question to answer, because part of it is keratinized (and therefore indicates this is skin) and part of it isn't (indicating conj). Could this be a junction between lid skin and conj? No, because no such junction exists (without intervening landmarks of the lid margin). So this is either skin that lost keratinization or conj that acquired it. But which is it?

The answer is clear once we compare the epithelia on the two sides of the change in keratinization status. The epithelium on the keratinized side is vastly thicker than that on the nonkeratinized side, strongly suggesting it (the keratinized epithelium) is neoplastic. Further, there is a sharp demarcation between the two areas, suggestive of a border between normal and neoplastic epithelium.



At first blush this seems an impossible question to answer, because part of it is keratinized (and therefore indicates this is skin) and part of it isn't (indicating conj). Could this be a junction between lid skin and conj? No, because no such junction exists (without intervening landmarks of the lid margin). So this is either skin that lost keratinization or conj that acquired it. But which is it?

The answer is clear once we compare the epithelia on the two sides of the change in keratinization status. The epithelium on the keratinized side is vastly thicker than that on the nonkeratinized side, strongly suggesting it (the keratinized epithelium) is neoplastic. Further, there is a sharp demarcation between the two areas, suggestive of a border between normal and neoplastic epithelium. Conj containing florid neoplasia + keratinization is strongly suggestive of

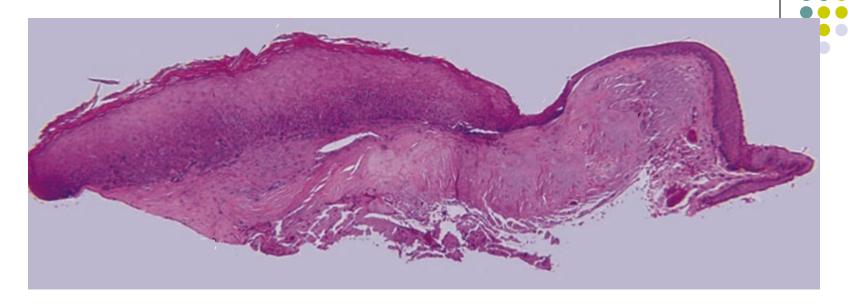


Again, first things first: Skin, or conj?

At first blush this seems an impossible question to answer, because part of it is keratinized (and therefore indicates this is skin) and part of it isn't (indicating conj). Could this be a junction between lid skin and conj? No, because no such junction exists (without intervening landmarks of the lid margin). So this is either skin that lost keratinization or conj that acquired it. But which is it?

The answer is clear once we compare the epithelia on the two sides of the change in keratinization status. The epithelium on the keratinized side is vastly thicker than that on the nonkeratinized side, strongly suggesting it (the keratinized epithelium) is neoplastic. Further, there is a sharp demarcation between the two areas, suggestive of a border between normal and neoplastic epithelium. Conj containing florid neoplasia + keratinization is strongly suggestive of ocular surface squamous neoplasia (OSSN).

146

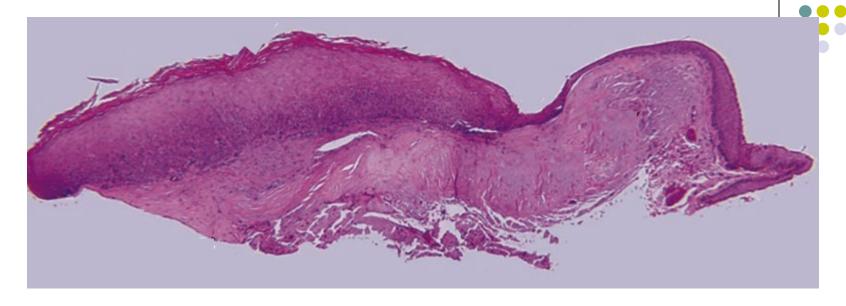


Again, first things first: Skin, or conj?

At first blush this seems an impossible question to answer, because part of it is keratinized (and therefore indicates this is skin) and part of it isn't (indicating conj). *Could this be a junction between lid skin and conj?* No, because no such junction exists (without intervening landmarks of the lid margin). So this is either skin that lost keratinization or conj that acquired it. But

OSSN arises on two-words portions of the conj

147

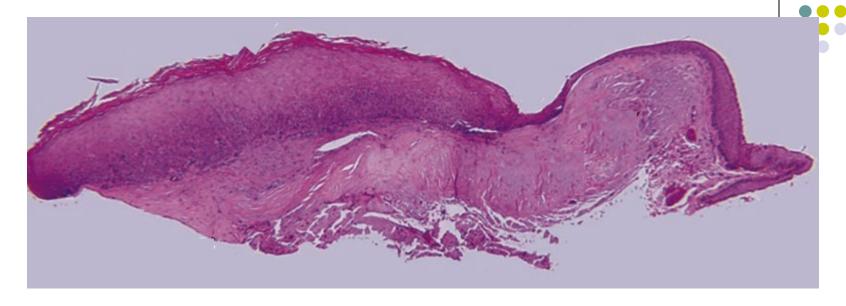


Again, first things first: Skin, or conj?

At first blush this seems an impossible question to answer, because part of it is keratinized (and therefore indicates this is skin) and part of it isn't (indicating conj). *Could this be a junction between lid skin and conj?* No, because no such junction exists (without intervening landmarks of the lid margin). So this is either skin that lost keratinization or conj that acquired it. But

OSSN arises on sun-exposed portions of the conj

148

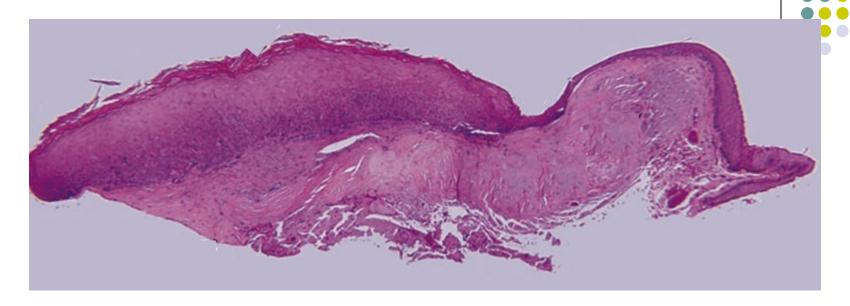


Again, first things first: Skin, or conj?

At first blush this seems an impossible question to answer, because part of it is keratinized (and therefore indicates this is skin) and part of it isn't (indicating conj). *Could this be a junction between lid skin and conj?* No, because no such junction exists (without intervening landmarks of the lid margin). So this is either skin that lost keratinization or conj that acquired it. But

OSSN arises on sun-exposed portions of the conj (sun exposure is a strong risk factor).

149

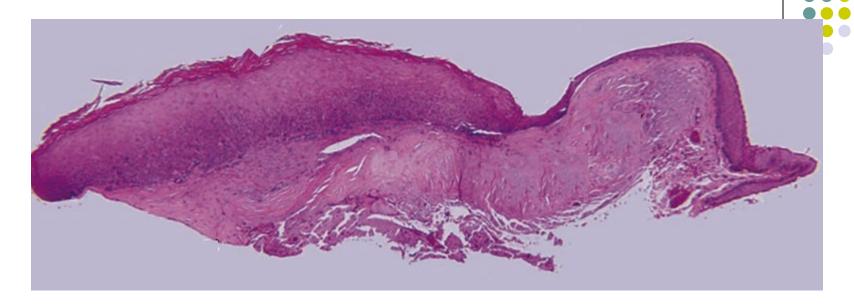


Again, first things first: Skin, or conj?

At first blush this seems an impossible question to answer, because part of it is keratinized (and therefore indicates this is skin) and part of it isn't (indicating conj). *Could this be a junction between lid skin and conj?* No, because no such junction exists (without intervening landmarks of the lid margin). So this is either skin that lost keratinization or conj that acquired it. But

OSSN arises on sun-exposed portions of the conj (sun exposure is a strong risk factor). It is more common in older vs journey individuals.

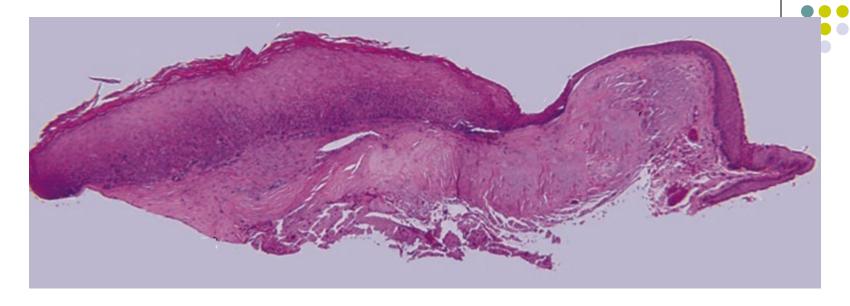
150



Again, first things first: Skin, or conj?

At first blush this seems an impossible question to answer, because part of it is keratinized (and therefore indicates this is skin) and part of it isn't (indicating conj). *Could this be a junction between lid skin and conj?* No, because no such junction exists (without intervening landmarks of the lid margin). So this is either skin that lost keratinization or conj that acquired it. But

OSSN arises on sun-exposed portions of the conj (sun exposure is a strong risk factor). It is more common in older individuals.

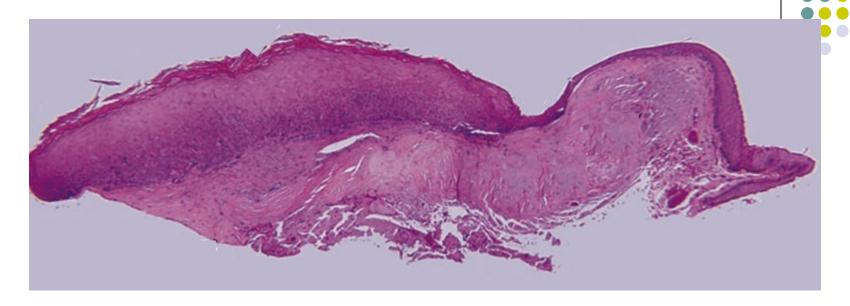


Again, first things first: Skin, or conj?

At first blush this seems an impossible question to answer, because part of it is keratinized (and therefore indicates this is skin) and part of it isn't (indicating conj). *Could this be a junction between lid skin and conj?* No, because no such junction exists (without intervening landmarks of the lid margin). So this is either skin that lost keratinization or conj that acquired it. But

OSSN arises on sun-exposed portions of the conj (sun exposure is a strong risk factor). It is more common in older individuals. (OSSN in individuals <50 should raise suspicion for block infection.)

152

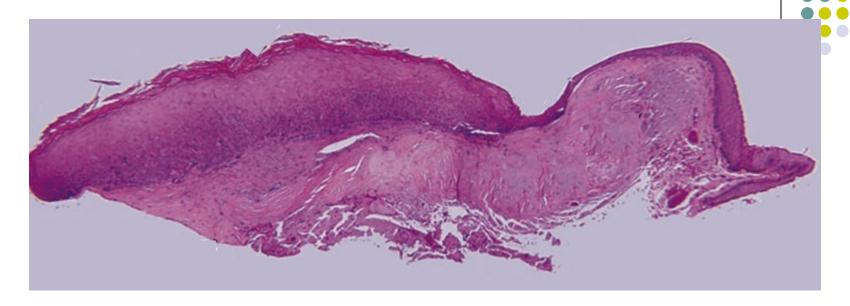


Again, first things first: Skin, or conj?

At first blush this seems an impossible question to answer, because part of it is keratinized (and therefore indicates this is skin) and part of it isn't (indicating conj). *Could this be a junction between lid skin and conj?* No, because no such junction exists (without intervening landmarks of the lid margin). So this is either skin that lost keratinization or conj that acquired it. But

OSSN arises on sun-exposed portions of the conj (sun exposure is a strong risk factor). It is more common in older individuals. (OSSN in individuals <50 should raise suspicion for HIV infection.)

153

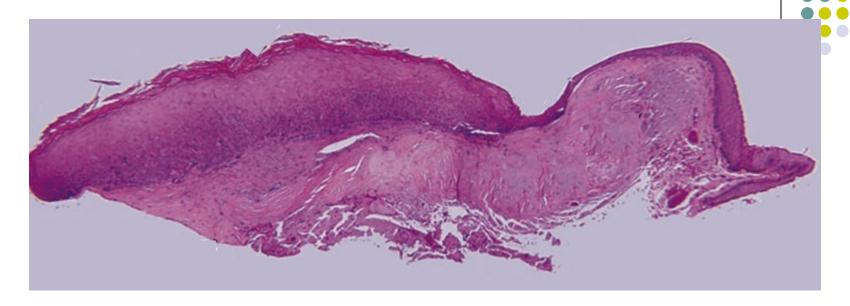


Again, first things first: Skin, or conj?

At first blush this seems an impossible question to answer, because part of it is keratinized (and therefore indicates this is skin) and part of it isn't (indicating conj). *Could this be a junction between lid skin and conj?* No, because no such junction exists (without intervening landmarks of the lid margin). So this is either skin that lost keratinization or conj that acquired it. But

OSSN arises on sun-exposed portions of the conj (sun exposure is a strong risk factor). It is more common in older individuals. (OSSN in individuals <50 should raise suspicion for HIV infection.) Keratinization pathognomonic for OSSN

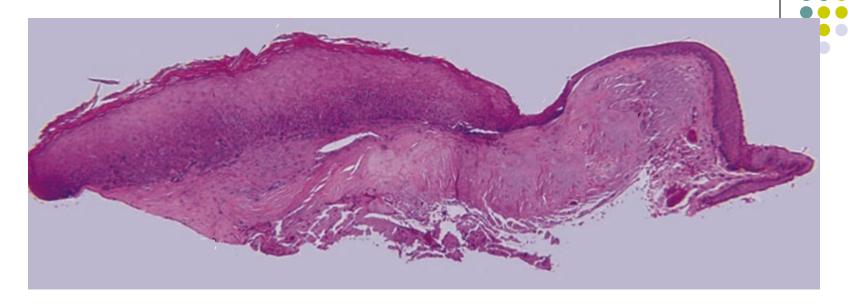
154



Again, first things first: Skin, or conj?

At first blush this seems an impossible question to answer, because part of it is keratinized (and therefore indicates this is skin) and part of it isn't (indicating conj). *Could this be a junction between lid skin and conj?* No, because no such junction exists (without intervening landmarks of the lid margin). So this is either skin that lost keratinization or conj that acquired it. But

OSSN arises on sun-exposed portions of the conj (sun exposure is a strong risk factor). It is more common in older individuals. (OSSN in individuals <50 should raise suspicion for HIV infection.) Keratinization isn't pathognomonic for OSSN

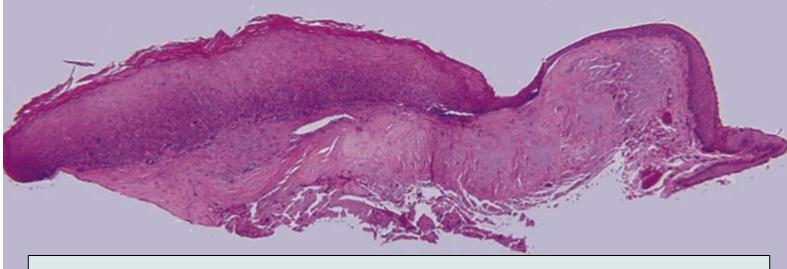


Again, first things first: Skin, or conj?

At first blush this seems an impossible question to answer, because part of it is keratinized (and therefore indicates this is skin) and part of it isn't (indicating conj). *Could this be a junction between lid skin and conj?* No, because no such junction exists (without intervening landmarks of the lid margin). So this is either skin that lost keratinization or conj that acquired it. But

OSSN arises on sun-exposed portions of the conj (sun exposure is a strong risk factor). It is more common in older individuals. (OSSN in individuals <50 should raise suspicion for HIV infection.) Keratinization isn't pathognomonic for OSSN; however, it is very common in it, and thus conj keratinization should prompt strong consideration of OSSN.

156



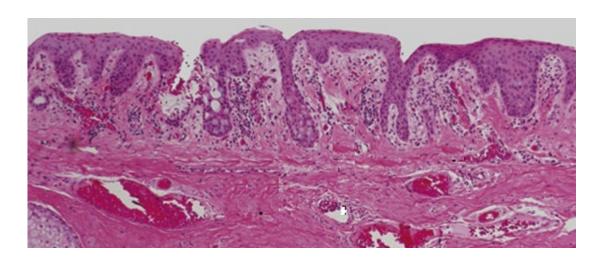
For more on OSSN, see slide-set K25

At first blush this seems an impossible question to answer, because part of it is keratinized (and therefore indicates this is skin) and part of it isn't (indicating conj). *Could this be a junction between lid skin and conj?* No, because no such junction exists (without intervening landmarks of the lid margin). So this is either skin that lost keratinization or conj that acquired it. But

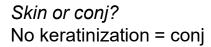
OSSN arises on sun-exposed portions of the conj (sun exposure is a strong risk factor). It is more common in older individuals. (OSSN in individuals <50 should raise suspicion for HIV infection.) Keratinization isn't pathognomonic for OSSN; however, it is very common in it, and thus conj keratinization should prompt strong consideration of OSSN.

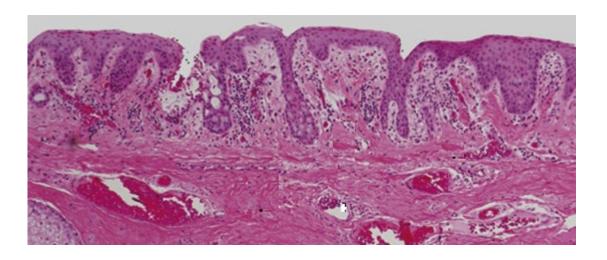


Skin or conj?

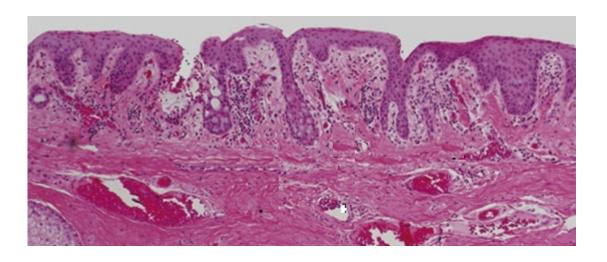








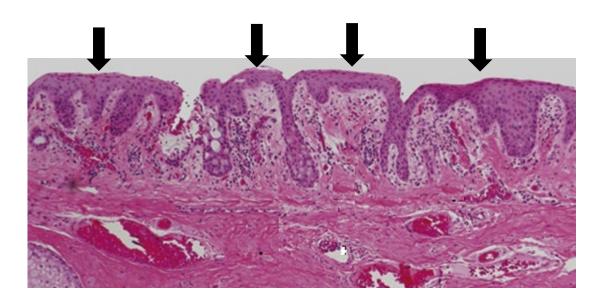




Skin or conj?
No keratinization = conj

There's a classic field mark here—what is it?



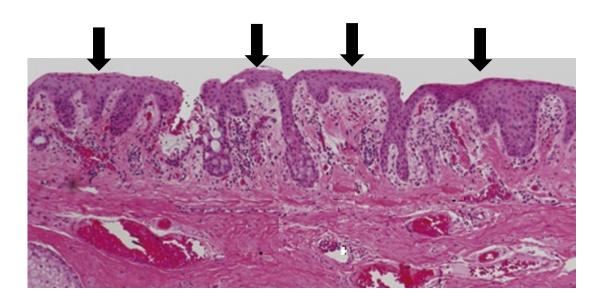


Skin or conj?
No keratinization = conj

There's a classic field mark here—what is it?

It's these closely packed, flat-topped mesa-looking structures. These correspond to a slit-lamp finding of



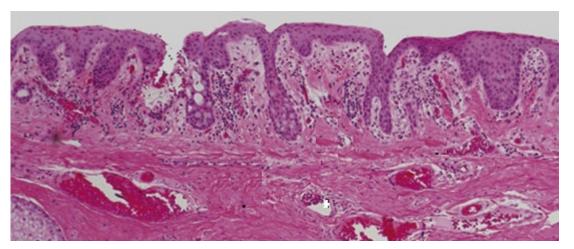


Skin or conj?
No keratinization = conj

There's a classic field mark here—what is it?

It's these closely packed, flat-topped mesa-looking structures. These correspond to a slit-lamp finding of 'cobblestones'.





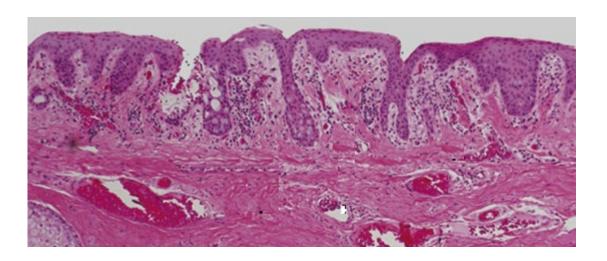
What's the diagnosis?

Skin or conj?
No keratinization = conj

There's a classic field mark here—what is it?

It's these closely packed, flat-topped mesa-looking structures. These correspond to a slit-lamp finding of 'cobblestones'. When you see these, one thing should come to mind:





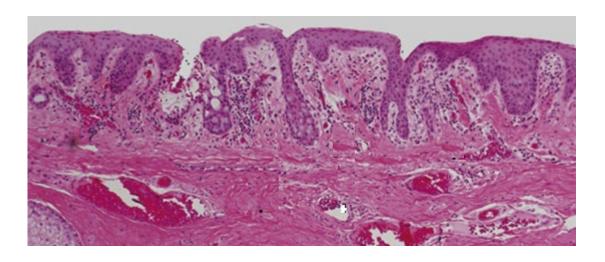
Skin or conj?
No keratinization = conj

There's a classic field mark here—what is it?

It's these closely packed, flat-topped mesa-looking structures. These correspond to a slit-lamp finding of 'cobblestones'. When you see these, one thing should come to mind:

Papillary conjunctivitis is one of # main forms of conjunctivitis





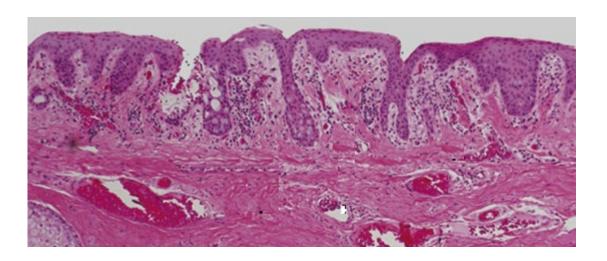
Skin or conj?
No keratinization = conj

There's a classic field mark here—what is it?

It's these closely packed, flat-topped mesa-looking structures. These correspond to a slit-lamp finding of 'cobblestones'. When you see these, one thing should come to mind:

Papillary conjunctivitis is one of two main forms of conjunctivitis





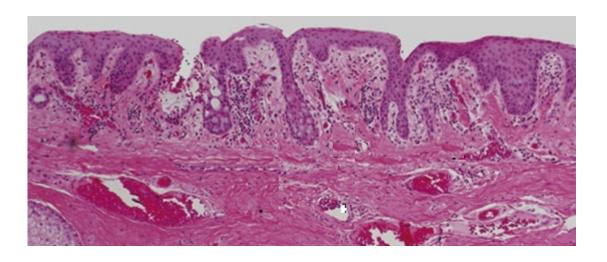
Skin or conj?
No keratinization = conj

There's a classic field mark here—what is it?

It's these closely packed, flat-topped mesa-looking structures. These correspond to a slit-lamp finding of 'cobblestones'. When you see these, one thing should come to mind:

Papillary conjunctivitis is one of two main forms of conjunctivitis (the other being





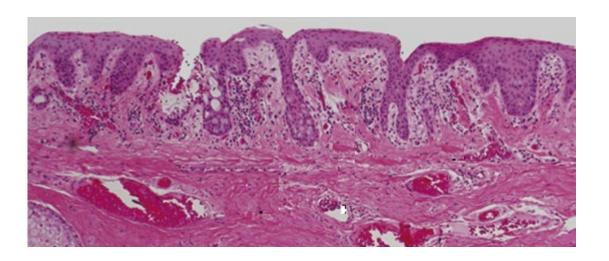
Skin or conj?
No keratinization = conj

There's a classic field mark here—what is it?

It's these closely packed, flat-topped mesa-looking structures. These correspond to a slit-lamp finding of 'cobblestones'. When you see these, one thing should come to mind:

Papillary conjunctivitis is one of two main forms of conjunctivitis (the other being follicular).





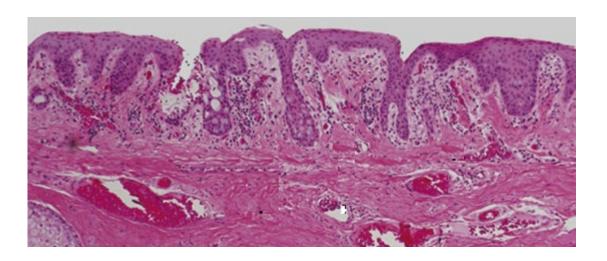
Skin or conj?
No keratinization = conj

There's a classic field mark here—what is it?

It's these closely packed, flat-topped mesa-looking structures. These correspond to a slit-lamp finding of 'cobblestones'. When you see these, one thing should come to mind:

Papillary conjunctivitis is one of two main forms of conjunctivitis (the other being *follicular*). It is most commonly associated with either broad dz cat. or a two words response.





Skin or conj?
No keratinization = conj

There's a classic field mark here—what is it?

It's these closely packed, flat-topped mesa-looking structures. These correspond to a slit-lamp finding of 'cobblestones'. When you see these, one thing should come to mind:

Papillary conjunctivitis is one of two main forms of conjunctivitis (the other being *follicular*). It is most commonly associated with either allergies or a foreign body response.



Skin or conj?
No keratinization = conj

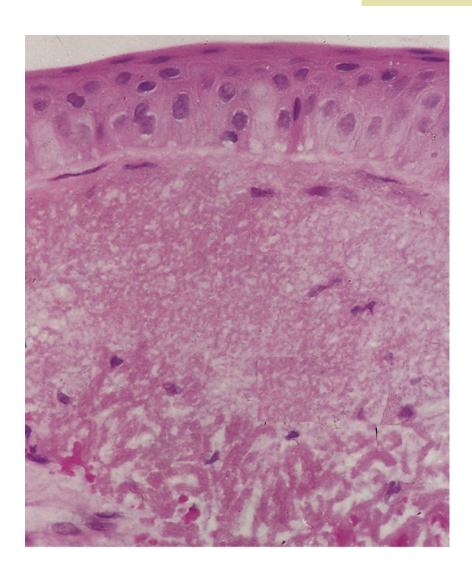
There's a classic field mark here—what is it?

It's these closely packed, flat-topped

These inding of see these, mind:

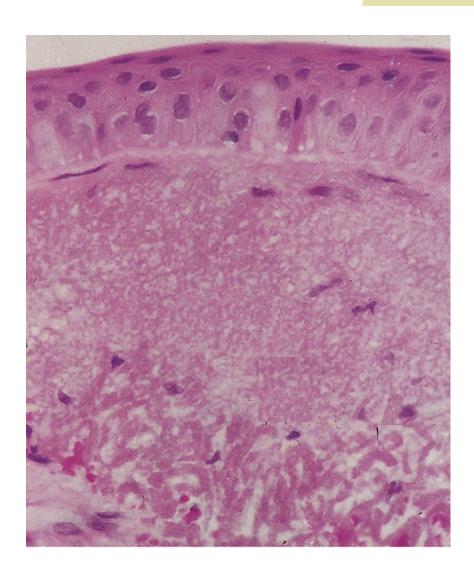
For more on papillary (and follicular) conjunctivitis, see slide-set K3

Papillary conjunctivitis is one of two main forms of conjunctivitis (the other being *follicular*). It is most commonly associated with either allergies or a foreign body response.



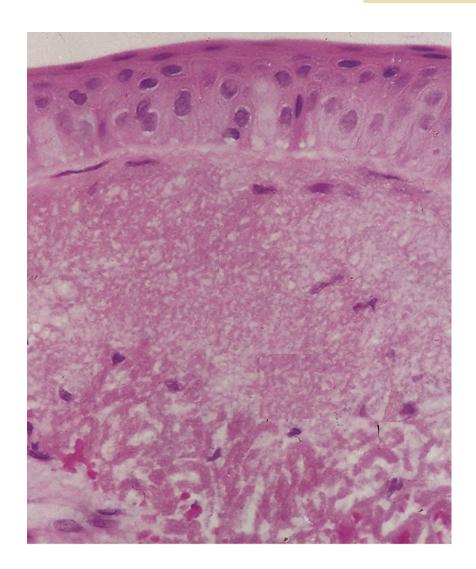
Skin or conj?





Skin or conj?
No keratinization = conj

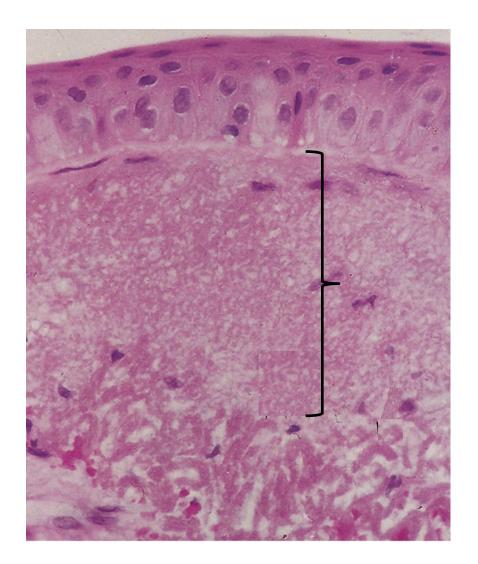






Skin or conj?
No keratinization = conj

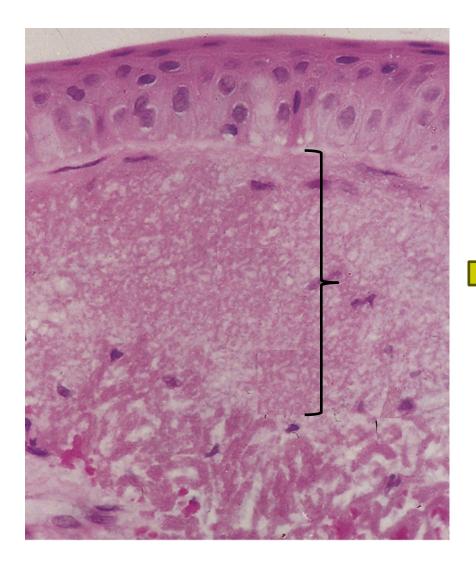
There's a good field mark here—what is it?





Skin or conj?
No keratinization = conj

There's a good field mark here—what is it? It's this large acellular region in the stroma.

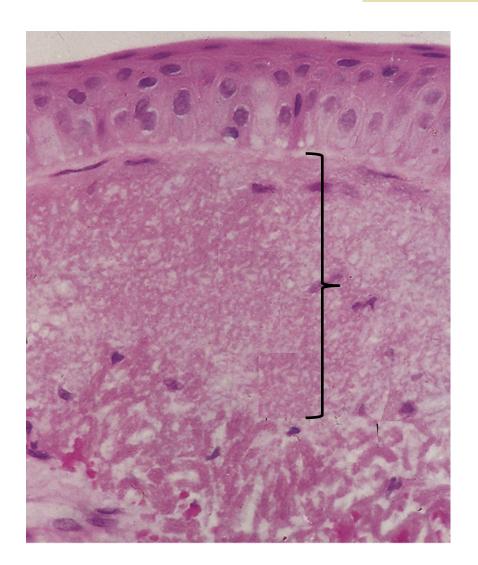


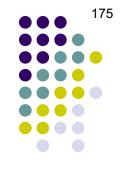


Skin or conj?
No keratinization = conj

There's a good field mark here—what is it? It's this large acellular region in the stroma. When you see this in a conj specimen, think

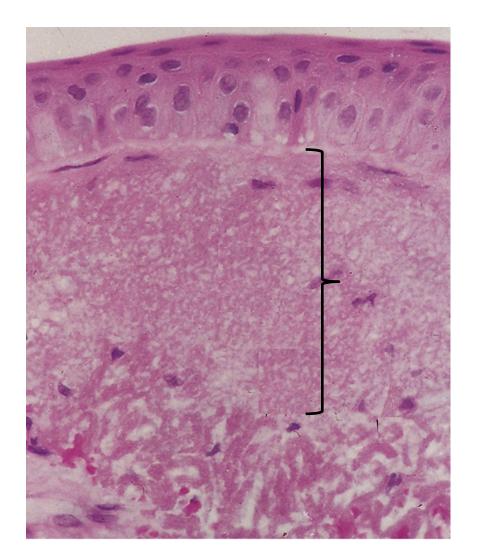
two words





Skin or conj?
No keratinization = conj

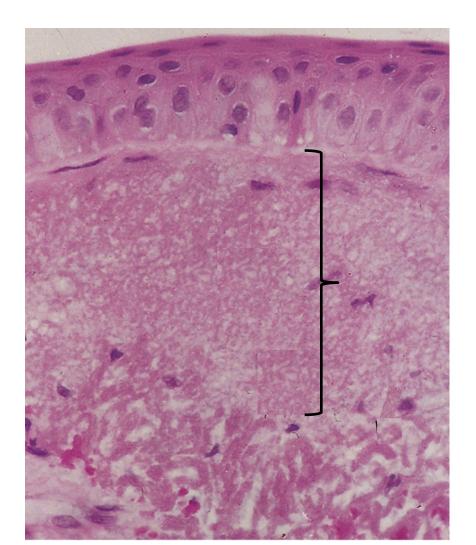
There's a good field mark here—what is it? It's this large acellular region in the stroma. When you see this in a conj specimen, think elastotic degeneration





Skin or conj?
No keratinization = conj

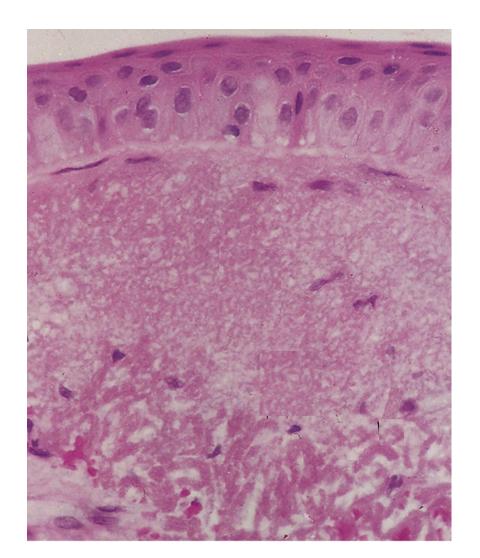
There's a good field mark here—what is it? It's this large acellular region in the stroma. When you see this in a conj specimen, think elastotic degeneration, which refers to fragmentation of stromal ______.





Skin or conj?
No keratinization = conj

There's a good field mark here—what is it? It's this large acellular region in the stroma. When you see this in a conj specimen, think elastotic degeneration, which refers to fragmentation of stromal collagen.

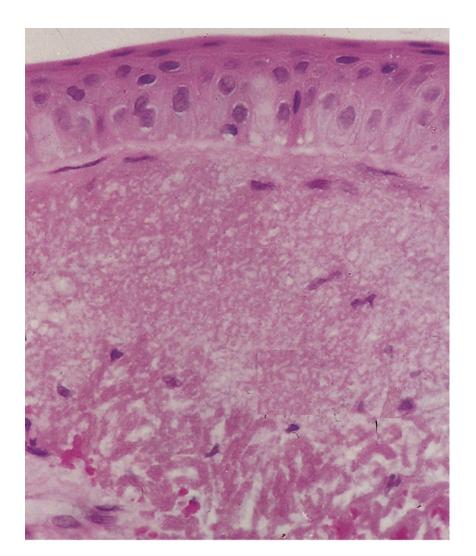




Skin or conj?
No keratinization = conj

There's a good field mark here—what is it?
It's this large acellular region in the stroma.
When you see this in a conj specimen, think elastotic degeneration, which refers to fragmentation of stromal collagen. If elastotic degeneration is present, it means you're looking at one of two related conditions:

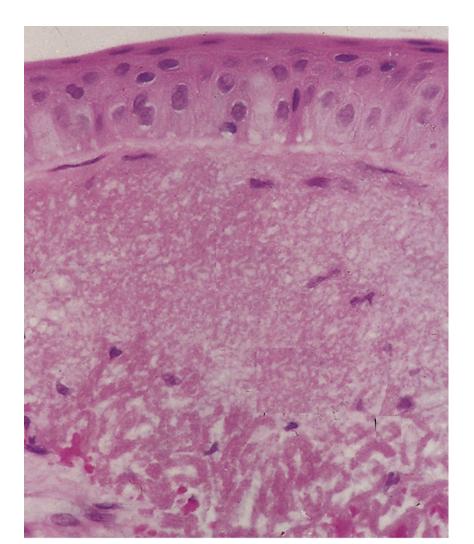
and





Skin or conj?
No keratinization = conj

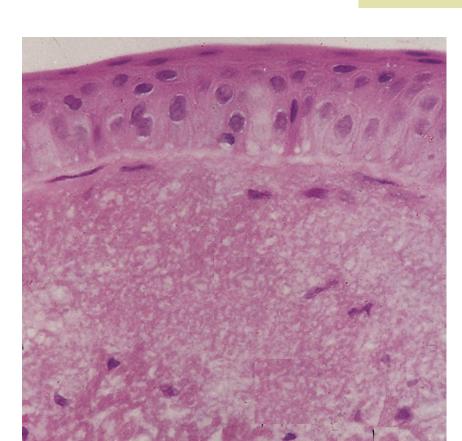
There's a good field mark here—what is it? It's this large acellular region in the stroma. When you see this in a conj specimen, think elastotic degeneration, which refers to fragmentation of stromal collagen. If elastotic degeneration is present, it means you're looking at one of two related conditions: Pinguecula and pterygium.





Skin or conj?
No keratinization = conj

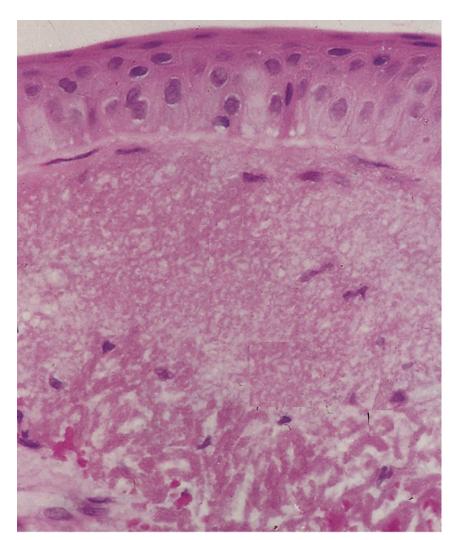
There's a good field mark here—what is it?
It's this large acellular region in the stroma.
When you see this in a conj specimen, think elastotic degeneration, which refers to fragmentation of stromal collagen. If elastotic degeneration is present, it means you're looking at one of two related conditions: Pinguecula and pterygium. These are distinguishable via whether prominent wowords are present (= pterygium) or absent (= pinguecula).





Skin or conj?
No keratinization = conj

There's a good field mark here—what is it? It's this large acellular region in the stroma. When you see this in a conj specimen, think elastotic degeneration, which refers to fragmentation of stromal collagen. If elastotic degeneration is present, it means you're looking at one of two related conditions: Pinguecula and pterygium. These are distinguishable via whether prominent blood vessels are present (= pterygium) or absent (= pinguecula).



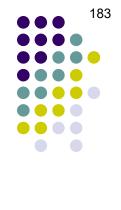
What's the diagnosis?



Skin or conj?
No keratinization = conj

There's a good field mark here—what is it?
It's this large acellular region in the stroma.
When you see this in a conj specimen, think
elastotic degeneration, which refers to
fragmentation of stromal collagen. If elastotic
degeneration is present, it means you're looking
at one of two related conditions: Pinguecula
and pterygium. These are distinguishable via
whether prominent blood vessels are present
(= pterygium) or absent (= pinguecula).

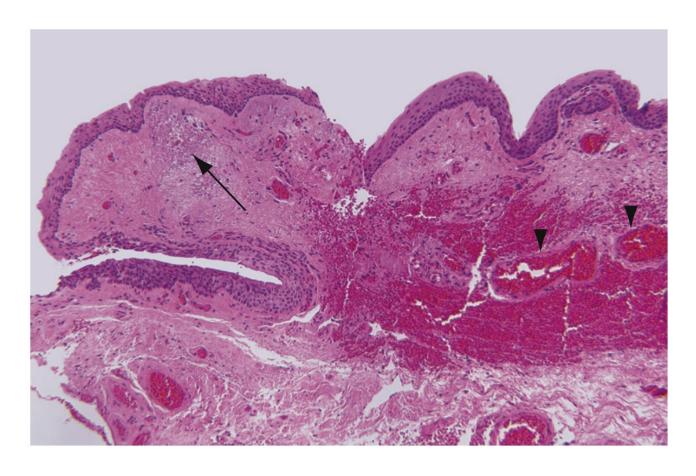




There's a good field mark here—what is it?
It's this large acellular region in the stroma.
When you see this in a conj specimen, think elastotic degeneration, which refers to fragmentation of stromal collagen. If elastotic degeneration is present, it means you're looking at one of two related conditions: Pinguecula and pterygium. These are distinguishable via whether prominent blood vessels are present (= pterygium) or absent (= pinguecula).

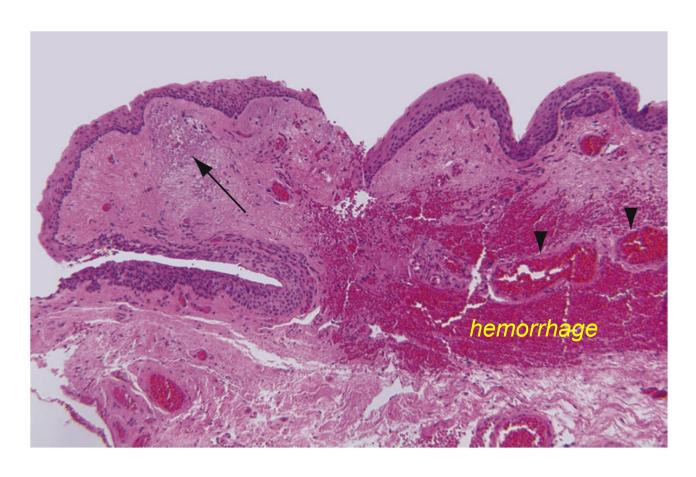
Pinguecula (no prominent blood vessels present)





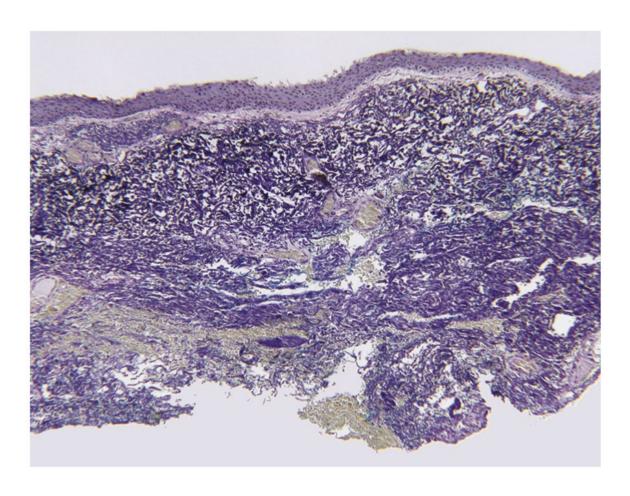
For comparison, here is a **pterygium**. Note the elastotic degeneration (*arrow*) as well as the blood vessels (*arrowheads*).





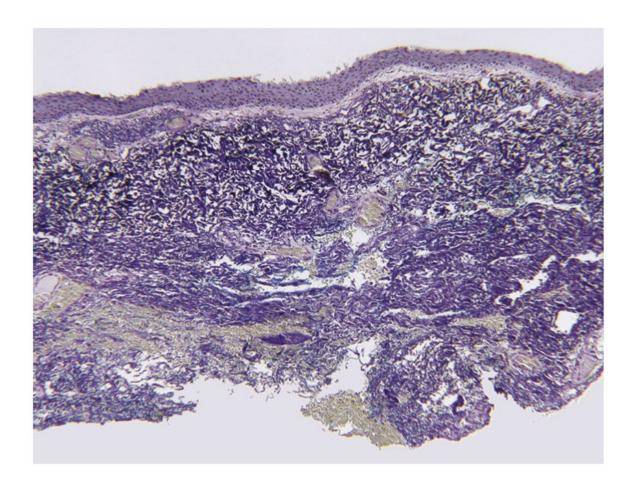
For comparison, here is a **pterygium**. Note the elastotic degeneration (*arrow*) as well as the blood vessels (*arrowheads*). Surgically-induced hemorrhage is present as well.





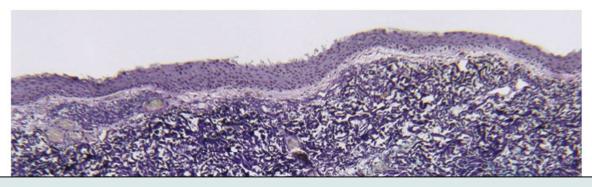
Protip: If a slide is stained for and is positive as all get-out like this, it's elastotic degeneration (and therefore a pinguecula or pterygium)



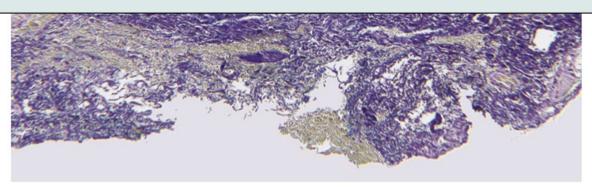


Protip: If a slide is stained for elastin and is positive as all get-out like this, it's elastotic degeneration (and therefore a pinguecula or pterygium)



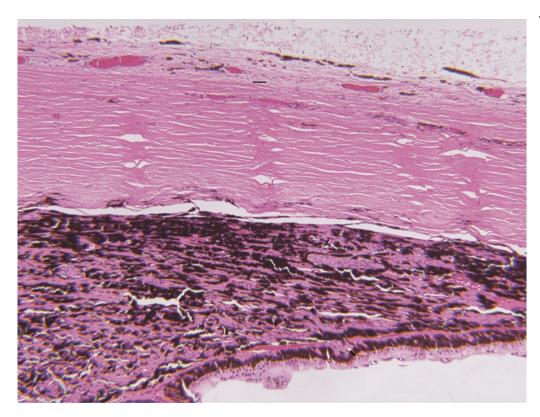


For more on pinguecula and pterygium, see slide-set K24



Protip: If a slide is stained for elastin and is positive as all get-out like this, it's elastotic degeneration (and therefore a pinguecula or pterygium)

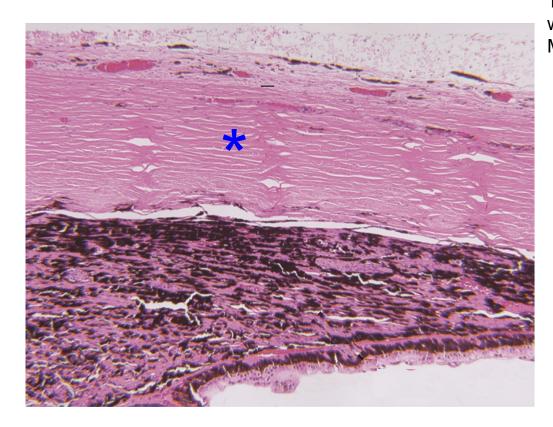




This one is puzzling at first. What tissue(s) are we looking at?

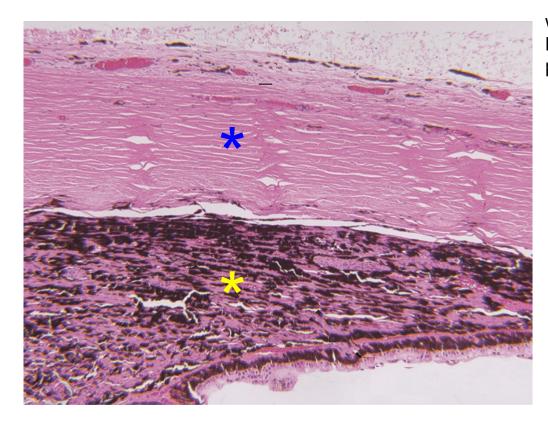
(Rhetorical question—keep going)





This one is puzzling at first. What tissue(s) are we looking at?
Maybe this is the cornea

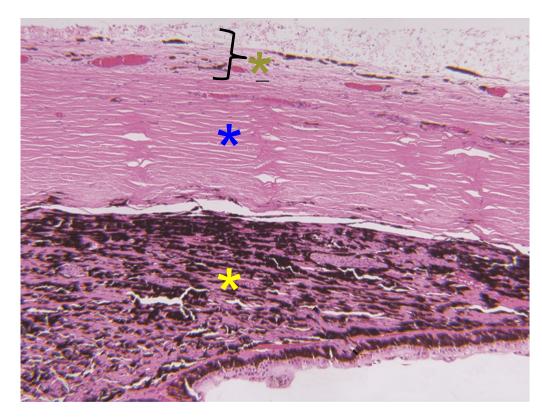




This one is puzzling at first. What tissue(s) are we looking at?

Maybe **this** is the cornea, and **this** is the iris pulled up against it in an angle closure event?

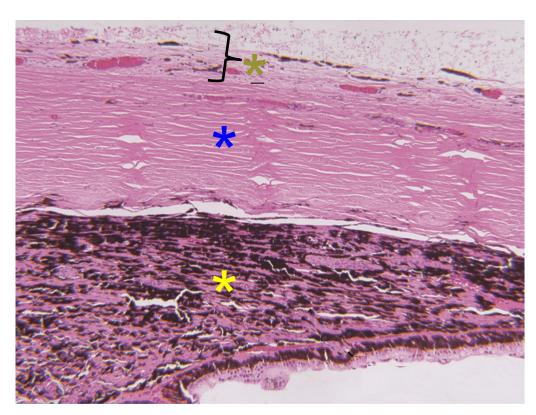




This one is puzzling at first. What tissue(s) are we looking at?

Maybe **this** is the cornea, and **this** is the iris pulled up against it in an angle closure event? Doesn't work, because it doesn't explain **this** tissue.





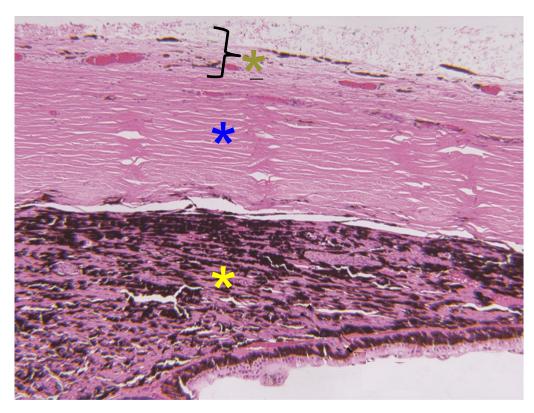
This one is puzzling at first. What tissue(s) are we looking at?

Maybe **this** is the cornea, and **this** is the iris pulled up against it in an angle closure event? Doesn't work, because it doesn't explain **this** tissue.

If **this** isn't the cornea, there's only one thing it could be—

. (Only the cornea and word have this laminar appearance.)



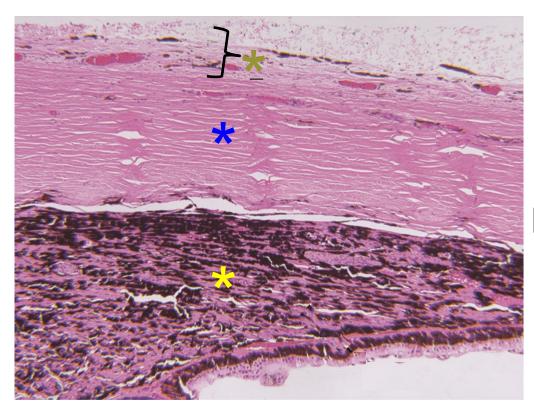


This one is puzzling at first. What tissue(s) are we looking at?

Maybe **this** is the cornea, and **this** is the iris pulled up against it in an angle closure event? Doesn't work, because it doesn't explain **this** tissue.

If **this** isn't the cornea, there's only one thing it could be—**sclera**. (Only the cornea and sclera have this laminar appearance.)



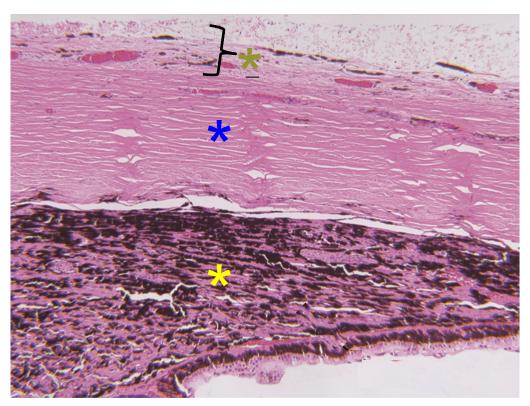


This one is puzzling at first. What tissue(s) are we looking at?

Maybe **this** is the cornea, and **this** is the iris pulled up against it in an angle closure event? Doesn't work, because it doesn't explain **this** tissue.

If **this** isn't the cornea, there's only one thing it could be—**sclera**. (Only the cornea and sclera have this laminar appearance.) And if **this** is sclera, this deeply pigmented tissue must be



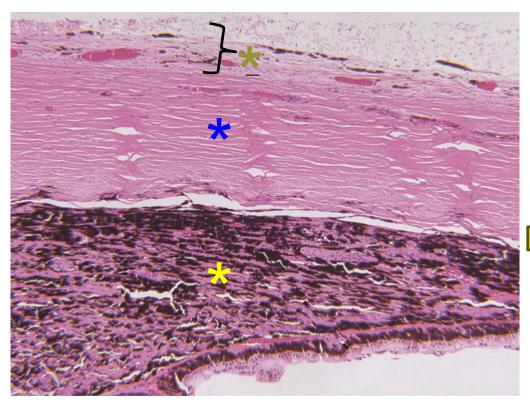


This one is puzzling at first. What tissue(s) are we looking at?

Maybe **this** is the cornea, and **this** is the iris pulled up against it in an angle closure event? Doesn't work, because it doesn't explain **this** tissue.

If **this** isn't the cornea, there's only one thing it could be—**sclera**. (Only the cornea and sclera have this laminar appearance.) And if **this** is sclera, this deeply pigmented tissue must be tree.



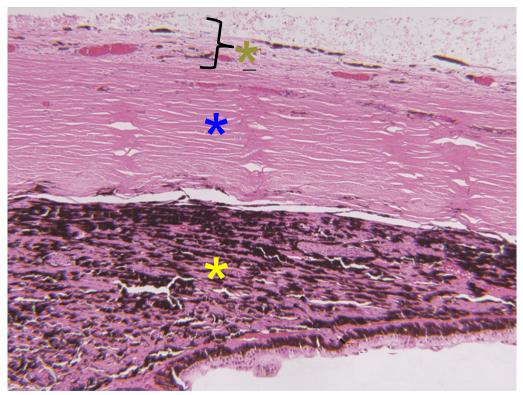


This one is puzzling at first. What tissue(s) are we looking at?

Maybe **this** is the cornea, and **this** is the iris pulled up against it in an angle closure event? Doesn't work, because it doesn't explain **this** tissue.

If **this** isn't the cornea, there's only one thing it could be—**sclera**. (Only the cornea and sclera have this laminar appearance.) And if **this** is sclera, this deeply pigmented tissue must be well. Which means **this** tissue must be



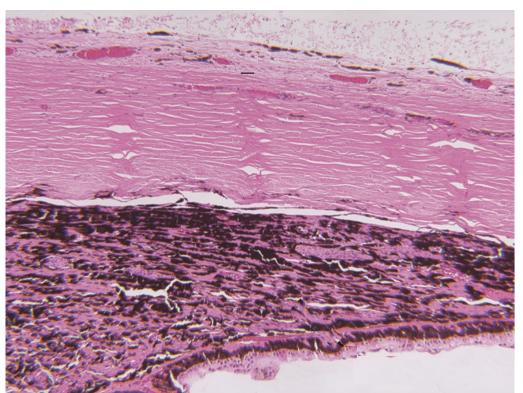


This one is puzzling at first. What tissue(s) are we looking at?

Maybe **this** is the cornea, and **this** is the iris pulled up against it in an angle closure event? Doesn't work, because it doesn't explain **this** tissue.

If **this** isn't the cornea, there's only one thing it could be—**sclera**. (Only the cornea and sclera have this laminar appearance.) And if **this** is sclera, this deeply pigmented tissue must be twee. Which means **this** tissue must be episclera.





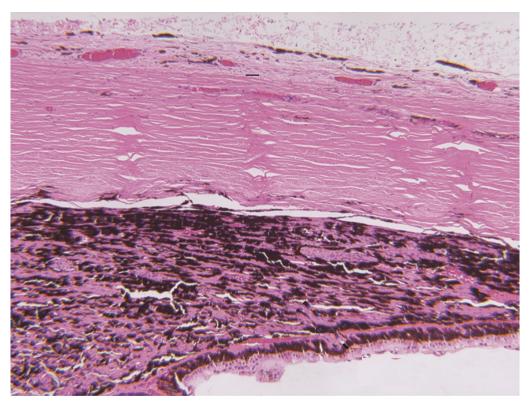
This one is puzzling at first. What tissue(s) are we looking at?

Maybe **this** is the cornea, and **this** is the iris pulled up against it in an angle closure event? Doesn't work, because it doesn't explain **this** tissue.

If **this** isn't the cornea, there's only one thing it could be—**sclera**. (Only the cornea and sclera have this laminar appearance.) And if **this** is sclera, this deeply pigmented tissue must be well. Which means **this** tissue must be episclera.

OK, now that we know where we are, what are we looking at, ie, what's the pathology? Look carefully at the sclera and (especially) the episclera—there's something unusual there. It's the relatively heavy presence of





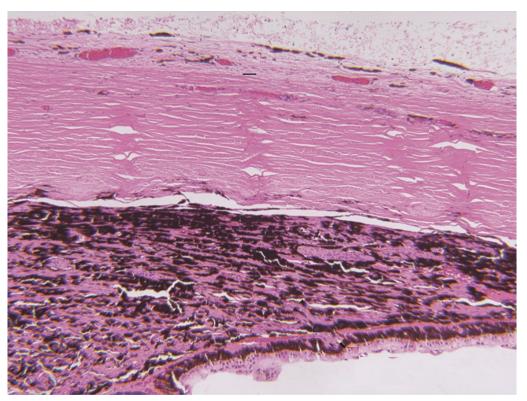
This one is puzzling at first. What tissue(s) are we looking at?

Maybe **this** is the cornea, and **this** is the iris pulled up against it in an angle closure event? Doesn't work, because it doesn't explain **this** tissue.

If **this** isn't the cornea, there's only one thing it could be—**sclera**. (Only the cornea and sclera have this laminar appearance.) And if **this** is sclera, this deeply pigmented tissue must be well. Which means **this** tissue must be episclera.

OK, now that we know where we are, what are we looking at, ie, what's the pathology? Look carefully at the sclera and (especially) the episclera—there's something unusual there. It's the relatively heavy presence of melanin/melanocytes.





What's the diagnosis?

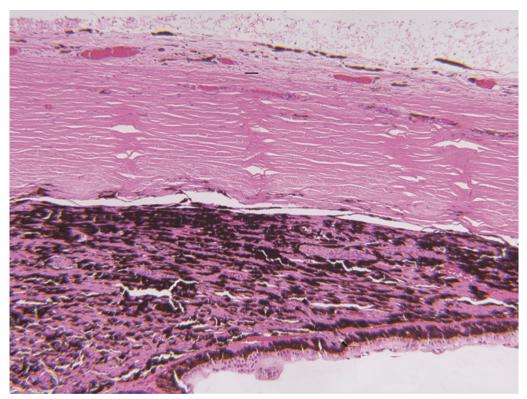
This one is puzzling at first. What tissue(s) are we looking at?

Maybe **this** is the cornea, and **this** is the iris pulled up against it in an angle closure event? Doesn't work, because it doesn't explain **this** tissue.

If **this** isn't the cornea, there's only one thing it could be—**sclera**. (Only the cornea and sclera have this laminar appearance.) And if **this** is sclera, this deeply pigmented tissue must be well. Which means **this** tissue must be episclera.

OK, now that we know where we are, what are we looking at, ie, what's the pathology? Look carefully at the sclera and (especially) the episclera—there's something unusual there. It's the relatively heavy presence of melanin/melanocytes. When you encounter heavy episcleral melanin-related findings, one thing should come to mind:





This one is puzzling at first. What tissue(s) are we looking at?

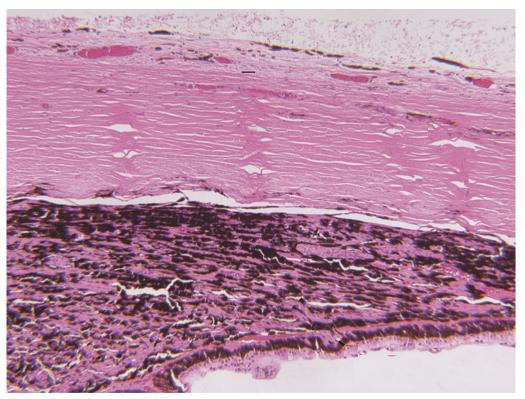
Maybe **this** is the cornea, and **this** is the iris pulled up against it in an angle closure event? Doesn't work, because it doesn't explain **this** tissue.

If **this** isn't the cornea, there's only one thing it could be—**sclera**. (Only the cornea and sclera have this laminar appearance.) And if this is sclera, **this** deeply pigmented tissue must be **uvea**. Which means **this** tissue must be **episclera**.

OK, now that we know where we are, what are we looking at, ie, what's the pathology? Look carefully at the sclera and (especially) the episclera—there's something unusual there. It's the relatively heavy presence of melanin/melanocytes. When you encounter heavy episcleral melanin-related findings, one thing should come to mind:

Ocular melanocytosis is a involving the deep episclera and sclera.





This one is puzzling at first. What tissue(s) are we looking at?

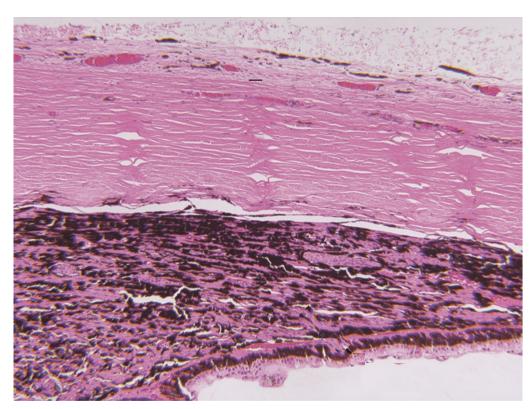
Maybe **this** is the cornea, and **this** is the iris pulled up against it in an angle closure event? Doesn't work, because it doesn't explain **this** tissue.

If **this** isn't the cornea, there's only one thing it could be—**sclera**. (Only the cornea and sclera have this laminar appearance.) And if this is sclera, **this** deeply pigmented tissue must be **uvea**. Which means **this** tissue must be **episclera**.

OK, now that we know where we are, what are we looking at, ie, what's the pathology? Look carefully at the sclera and (especially) the episclera—there's something unusual there. It's the relatively heavy presence of melanin/melanocytes. When you encounter heavy episcleral melanin-related findings, one thing should come to mind:

Ocular melanocytosis is a nevus involving the deep episclera and sclera.





This one is puzzling at first. What tissue(s) are we looking at?

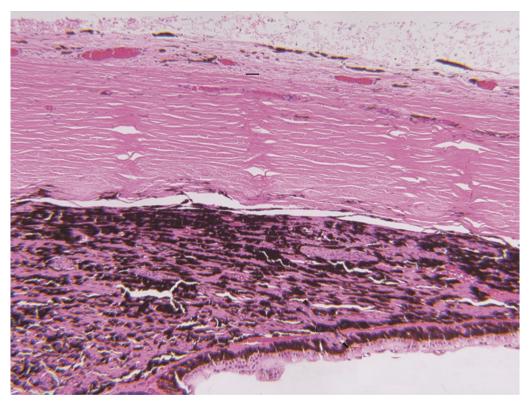
Maybe **this** is the cornea, and **this** is the iris pulled up against it in an angle closure event? Doesn't work, because it doesn't explain **this** tissue.

If **this** isn't the cornea, there's only one thing it could be—**sclera**. (Only the cornea and sclera have this laminar appearance.) And if this is sclera, **this** deeply pigmented tissue must be **uvea**. Which means **this** tissue must be **episclera**.

OK, now that we know where we are, what are we looking at, ie, what's the pathology? Look carefully at the sclera and (especially) the episclera—there's something unusual there. It's the relatively heavy presence of melanin/melanocytes. When you encounter heavy episcleral melanin-related findings, one thing should come to mind:

Ocular melanocytosis is a nevus involving the deep episclera and sclera. If the periocular skin is also involved, the condition is called melanocytosis





This one is puzzling at first. What tissue(s) are we looking at?

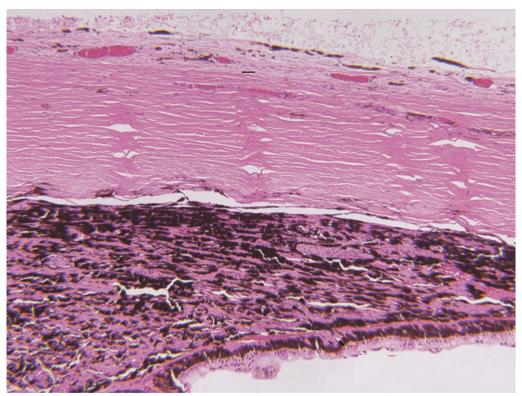
Maybe **this** is the cornea, and **this** is the iris pulled up against it in an angle closure event? Doesn't work, because it doesn't explain **this** tissue.

If **this** isn't the cornea, there's only one thing it could be—**sclera**. (Only the cornea and sclera have this laminar appearance.) And if this is sclera, **this** deeply pigmented tissue must be **uvea**. Which means **this** tissue must be **episclera**.

OK, now that we know where we are, what are we looking at, ie, what's the pathology? Look carefully at the sclera and (especially) the episclera—there's something unusual there. It's the relatively heavy presence of melanin/melanocytes. When you encounter heavy episcleral melanin-related findings, one thing should come to mind:

Ocular melanocytosis is a nevus involving the deep episclera and sclera. If the periocular skin is also involved, the condition is called oculodermal melanocytosis





This one is puzzling at first. What tissue(s) are we looking at?

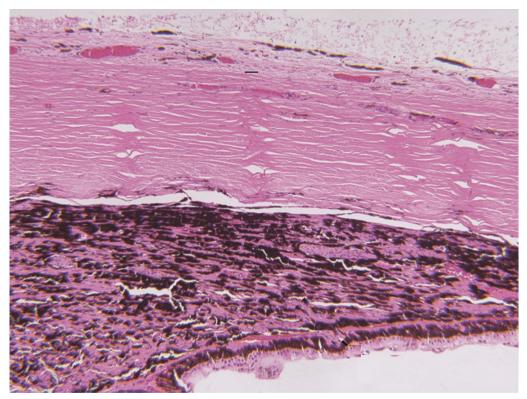
Maybe **this** is the cornea, and **this** is the iris pulled up against it in an angle closure event? Doesn't work, because it doesn't explain **this** tissue.

If **this** isn't the cornea, there's only one thing it could be—**sclera**. (Only the cornea and sclera have this laminar appearance.) And if this is sclera, **this** deeply pigmented tissue must be **uvea**. Which means **this** tissue must be **episclera**.

OK, now that we know where we are, what are we looking at, ie, what's the pathology? Look carefully at the sclera and (especially) the episclera—there's something unusual there. It's the relatively heavy presence of melanin/melanocytes. When you encounter heavy episcleral melanin-related findings, one thing should come to mind:

Ocular melanocytosis is a nevus involving the deep episclera and sclera. If the periocular skin is also involved, the condition is called oculodermal melanocytosis (aka three words).





This one is puzzling at first. What tissue(s) are we looking at?

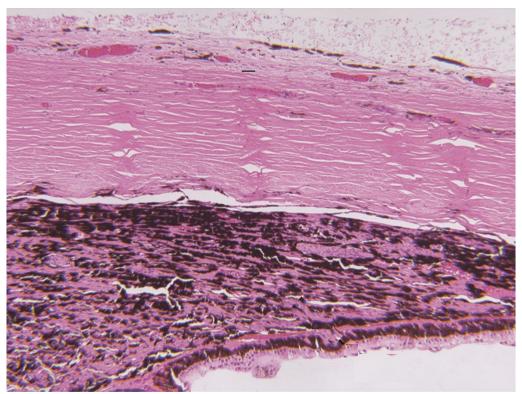
Maybe **this** is the cornea, and **this** is the iris pulled up against it in an angle closure event? Doesn't work, because it doesn't explain **this** tissue.

If **this** isn't the cornea, there's only one thing it could be—**sclera**. (Only the cornea and sclera have this laminar appearance.) And if this is sclera, **this** deeply pigmented tissue must be **uvea**. Which means **this** tissue must be **episclera**.

OK, now that we know where we are, what are we looking at, ie, what's the pathology? Look carefully at the sclera and (especially) the episclera—there's something unusual there. It's the relatively heavy presence of melanin/melanocytes. When you encounter heavy episcleral melanin-related findings, one thing should come to mind:

Ocular melanocytosis is a nevus involving the deep episclera and sclera. If the periocular skin is also involved, the condition is called oculodermal melanocytosis (aka Nevus of Ota).





This one is puzzling at first. What tissue(s) are we looking at?

Maybe **this** is the cornea, and **this** is the iris pulled up against it in an angle closure event? Doesn't work, because it doesn't explain **this** tissue.

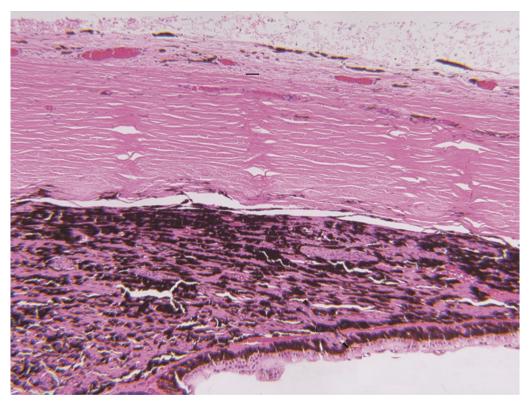
If **this** isn't the cornea, there's only one thing it could be—**sclera**. (Only the cornea and sclera have this laminar appearance.) And if this is sclera, **this** deeply pigmented tissue must be **uvea**. Which means **this** tissue must be **episclera**.

OK, now that we know where we are, what are we looking at, ie, what's the pathology? Look carefully at the sclera and (especially) the episclera—there's something unusual there. It's the relatively heavy presence of melanin/melanocytes. When you encounter heavy episcleral melanin-related findings, one thing should come to mind:

Ocular melanocytosis is a nevus involving the deep episclera and sclera. If the periocular skin is also involved, the condition is called oculodermal melanocytosis (aka *Nevus of Ota*).

Lightly vs pigmented individuals with melanocytosis are at significantly increased of





This one is puzzling at first. What tissue(s) are we looking at?

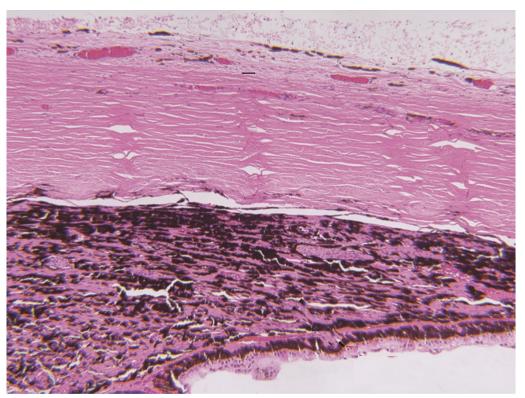
Maybe **this** is the cornea, and **this** is the iris pulled up against it in an angle closure event? Doesn't work, because it doesn't explain **this** tissue.

If **this** isn't the cornea, there's only one thing it could be—**sclera**. (Only the cornea and sclera have this laminar appearance.) And if this is sclera, **this** deeply pigmented tissue must be **uvea**. Which means **this** tissue must be **episclera**.

OK, now that we know where we are, what are we looking at, ie, what's the pathology? Look carefully at the sclera and (especially) the episclera—there's something unusual there. It's the relatively heavy presence of melanin/melanocytes. When you encounter heavy episcleral melanin-related findings, one thing should come to mind:

Ocular melanocytosis is a nevus involving the deep episclera and sclera. If the periocular skin is also involved, the condition is called oculodermal melanocytosis (aka *Nevus of Ota*). Lightly pigmented individuals with melanocytosis are at significantly increased of melanoma





This one is puzzling at first. What tissue(s) are we looking at?

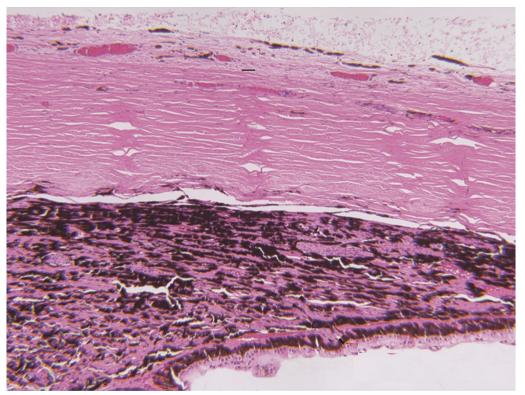
Maybe **this** is the cornea, and **this** is the iris pulled up against it in an angle closure event? Doesn't work, because it doesn't explain **this** tissue.

If **this** isn't the cornea, there's only one thing it could be—**sclera**. (Only the cornea and sclera have this laminar appearance.) And if this is sclera, **this** deeply pigmented tissue must be **uvea**. Which means **this** tissue must be **episclera**.

OK, now that we know where we are, what are we looking at, ie, what's the pathology? Look carefully at the sclera and (especially) the episclera—there's something unusual there. It's the relatively heavy presence of melanin/melanocytes. When you encounter heavy episcleral melanin-related findings, one thing should come to mind:

Ocular melanocytosis is a nevus involving the deep episclera and sclera. If the periocular skin is also involved, the condition is called oculodermal melanocytosis (aka *Nevus of Ota*). Lightly pigmented individuals with melanocytosis are at significantly increased of melanoma, almost always of the structure





This one is puzzling at first. What tissue(s) are we looking at?

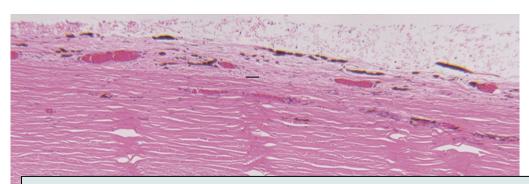
Maybe **this** is the cornea, and **this** is the iris pulled up against it in an angle closure event? Doesn't work, because it doesn't explain **this** tissue.

If **this** isn't the cornea, there's only one thing it could be—**sclera**. (Only the cornea and sclera have this laminar appearance.) And if this is sclera, **this** deeply pigmented tissue must be **uvea**. Which means **this** tissue must be **episclera**.

OK, now that we know where we are, what are we looking at, ie, what's the pathology? Look carefully at the sclera and (especially) the episclera—there's something unusual there. It's the relatively heavy presence of melanin/melanocytes. When you encounter heavy episcleral melanin-related findings, one thing should come to mind:

Ocular melanocytosis is a nevus involving the deep episclera and sclera. If the periocular skin is also involved, the condition is called oculodermal melanocytosis (aka *Nevus of Ota*). Lightly pigmented individuals with melanocytosis are at significantly increased of melanoma, almost always of the uvea.



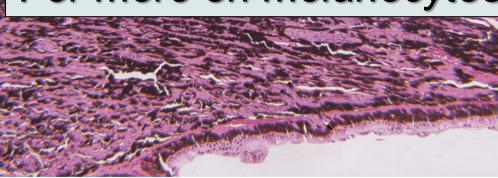


This one is puzzling at first. What tissue(s) are we looking at?

Maybe **this** is the cornea, and **this** is the iris pulled up against it in an angle closure event? Doesn't work, because it doesn't explain **this** tissue.

If **this** isn't the cornea, there's only one thing it could be—**sclera**. (Only the cornea and sclera have this laminar appearance.) And if this is

For more on melanocytosis, see slide-set O9

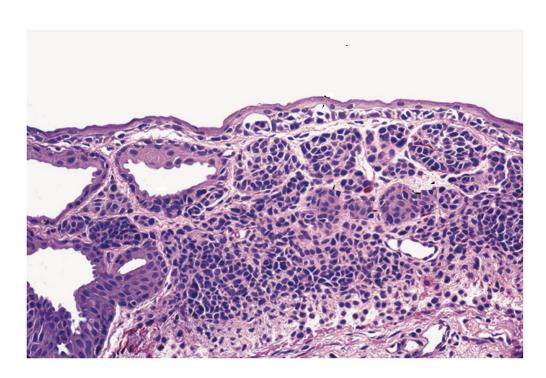


episciera .

OK, now that we know where we are, what are we looking at, ie, what's the pathology? Look carefully at the sclera and (especially) the episclera—there's something unusual there. It's the relatively heavy presence of melanin/melanocytes. When you encounter heavy episcleral melanin-related findings, one thing should come to mind:

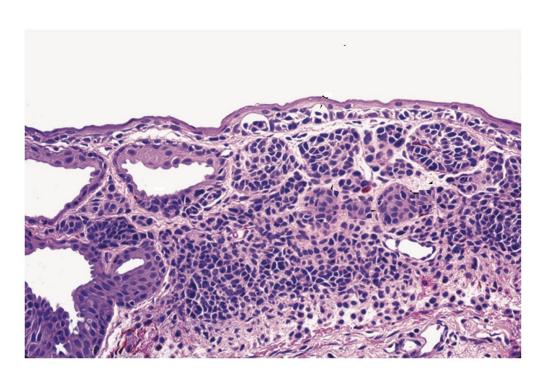
Ocular melanocytosis is a nevus involving the deep episclera and sclera. If the periocular skin is also involved, the condition is called oculodermal melanocytosis (aka *Nevus of Ota*). Lightly pigmented individuals with melanocytosis are at significantly increased of melanoma, almost always of the uvea.





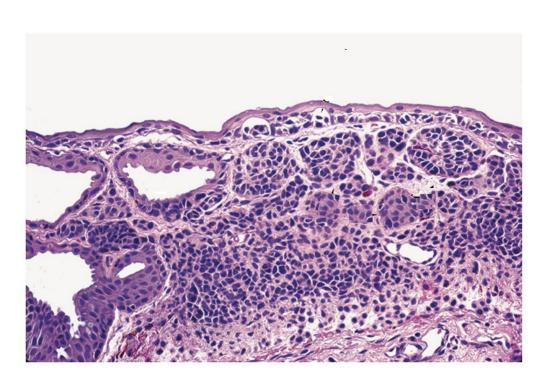
Skin/conj?





*Skin/conj?*Conj

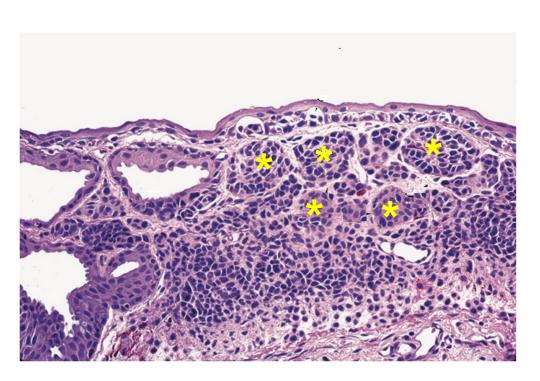




Skin/conj? Conj

There are a couple of field marks that, taken together, nail the diagnosis. What are they?





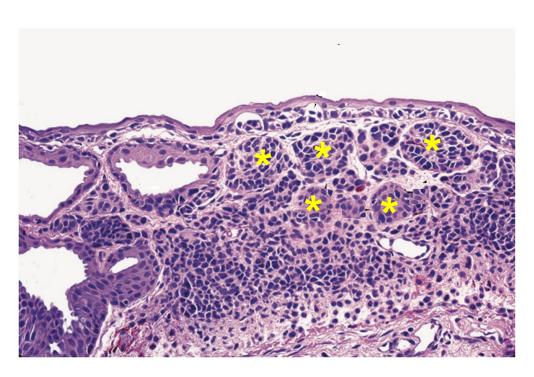
Skin/conj? Conj

There are a couple of field marks that, taken together, nail the diagnosis. What are they?

-- These tight clusters of cell type are called

--?





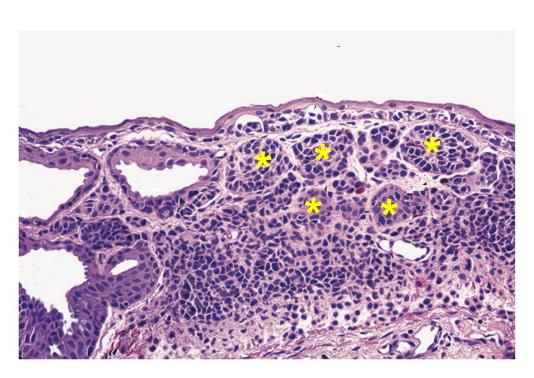
Skin/conj? Conj

There are a couple of field marks that, taken together, nail the diagnosis. What are they?

--These tight clusters of melanocytes are called nests

-- ?





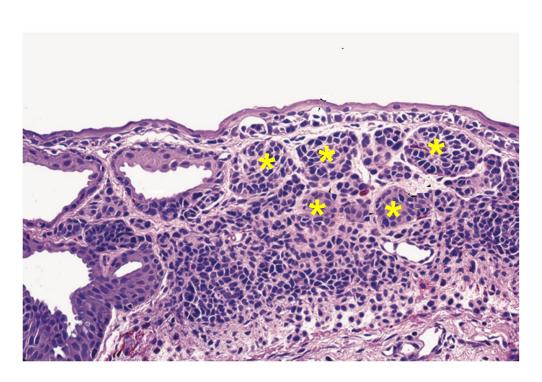
Skin/conj? Conj

There are a couple of field marks that, taken together, nail the diagnosis. What are they?

-- These tight clusters of melanocytes are called nests (aka

--?





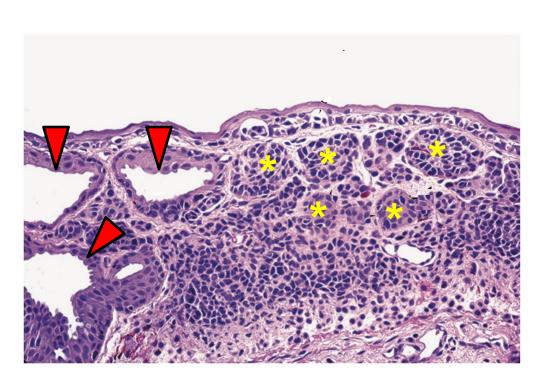
Skin/conj? Conj

There are a couple of field marks that, taken together, nail the diagnosis. What are they?

-- These tight clusters of melanocytes are called nests (aka theques)

--?





Skin/conj? Conj

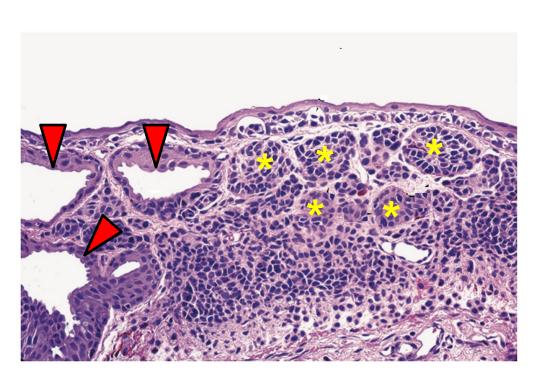
There are a couple of field marks that, taken together, nail the diagnosis. What are they?

-- These tight clusters of melanocytes are called nests (aka theques) and

--these

three words





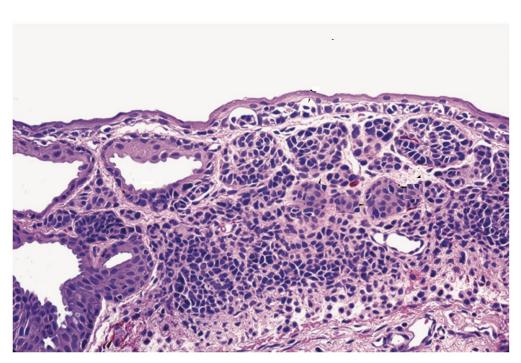
Skin/conj? Conj

There are a couple of field marks that, taken together, nail the diagnosis. What are they?

--These tight clusters of melanocytes are called nests (aka theques) and

--these epithelial inclusion cysts





What's the diagnosis?

Skin/conj? Conj

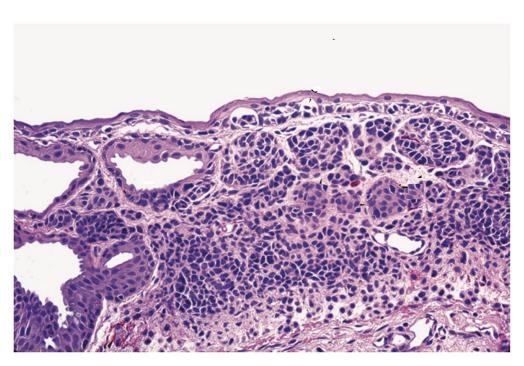
There are a couple of field marks that, taken together, nail the diagnosis. What are they?

--These tight clusters of melanocytes are called nests (aka theques) and

--these epithelial inclusion cysts

When you encounter melanocytes in nests associated with cysts, one condition should come to mind:





Skin/conj? Conj

There are a couple of field marks that, taken together, nail the diagnosis. What are they? -- These tight clusters of melanocytes are called nests (aka theques) and --these epithelial inclusion cysts

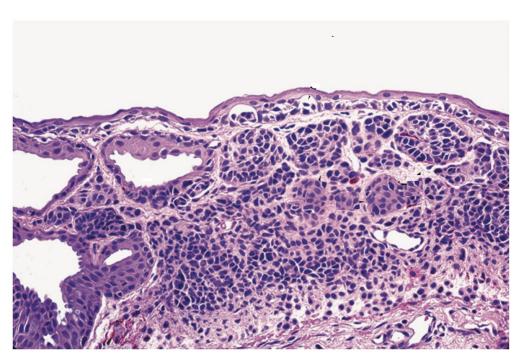
When you encounter melanocytes in nests associated with cysts, one condition should come to mind:

What's the diagnosis?

Melanocytic nevi almost always appear on the portion

conj





What's the diagnosis?

Skin/conj? Conj

There are a couple of field marks that, taken together, nail the diagnosis. What are they?

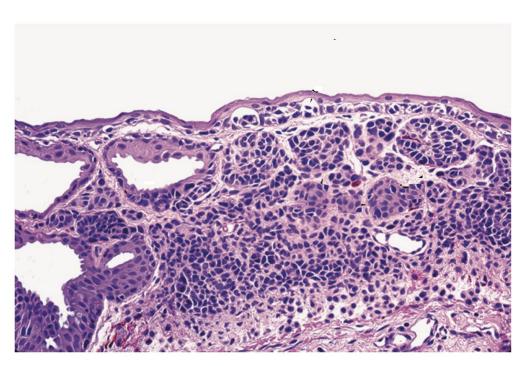
-- These tight clusters of melanocytes are called nests (aka theques) and

-- these epithelial inclusion cysts

When you encounter melanocytes in nests associated with cysts, one condition should come to mind:

Melanocytic nevi almost always appear on the bulbar conj





Skin/conj? Conj

There are a couple of field marks that, taken together, nail the diagnosis. What are they?

--These tight clusters of melanocytes are called nests (aka theques) and

--these epithelial inclusion cysts

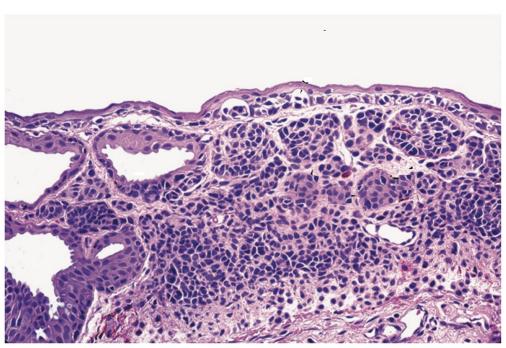
When you encounter melanocytes in nests associated with cysts, one condition should come to mind:

What's the diagnosis?

Melanocytic nevi almost always appear on the bulbar conj during

life stage





What's the diagnosis?

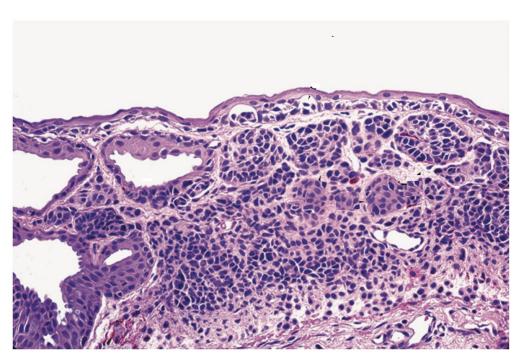
Skin/conj? Conj

There are a couple of field marks that, taken together, nail the diagnosis. What are they? -- These tight clusters of melanocytes are called nests (aka theques) and --these epithelial inclusion cysts

When you encounter melanocytes in nests associated with cysts, one condition should come to mind:

Melanocytic nevi almost always appear on the bulbar conj during childhood.





Skin/conj? Conj

There are a couple of field marks that, taken together, nail the diagnosis. What are they?

--These tight clusters of melanocytes are called nests (aka theques) and

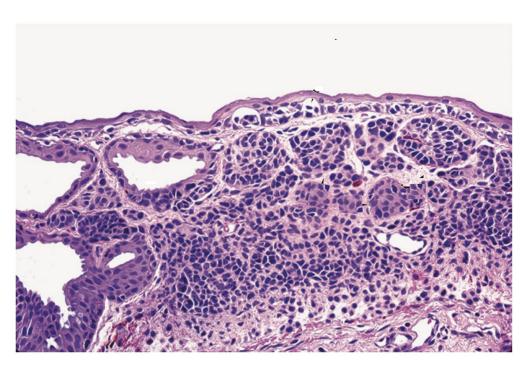
--these epithelial inclusion cysts

When you encounter melanocytes in nests associated with cysts, one condition should come to mind:

What's the diagnosis?

Melanocytic nevi almost always appear on the bulbar conj during childhood. As with cutaneous nevi, they evolve through a series of histologic conformations, from (in order) to





Skin/conj? Conj

There are a couple of field marks that, taken together, nail the diagnosis. What are they?

--These tight clusters of melanocytes are called nests (aka theques) and

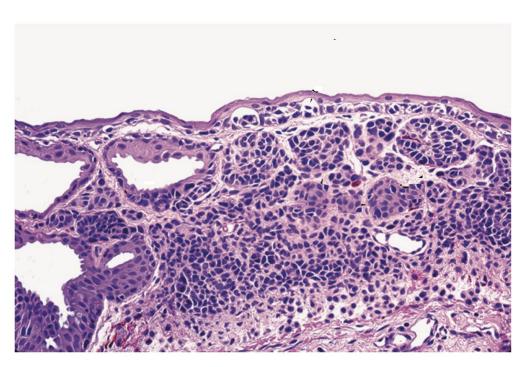
--these epithelial inclusion cysts

When you encounter melanocytes in nests associated with cysts, one condition should come to mind:

What's the diagnosis?

Melanocytic nevi almost always appear on the bulbar conj during childhood. As with cutaneous nevi, they evolve through a series of histologic conformations, from (in order) junctional to compound to stromal.





Skin/conj? Conj

There are a couple of field marks that, taken together, nail the diagnosis. What are they?

--These tight clusters of melanocytes are called nests (aka theques) and

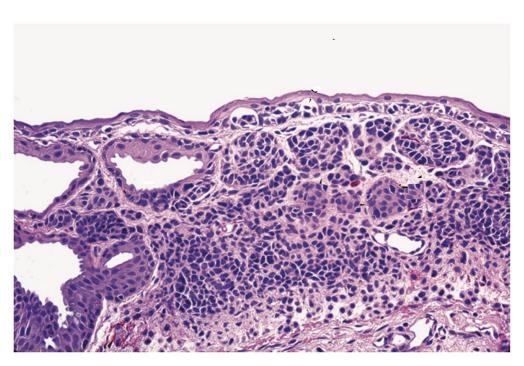
--these epithelial inclusion cysts

When you encounter melanocytes in nests associated with cysts, one condition should come to mind:

What's the diagnosis?

Melanocytic nevi almost always appear on the bulbar conj during childhood. As with cutaneous nevi, they evolve through a series of histologic conformations, from (in order) junctional to compound to stromal. Conj nevi have malignant potential.





Skin/conj? Conj

There are a couple of field marks that, taken together, nail the diagnosis. What are they?

--These tight clusters of melanocytes are called nests (aka theques) and

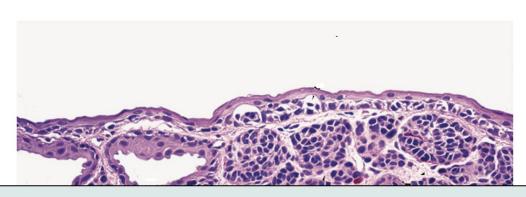
--these epithelial inclusion cysts

When you encounter melanocytes in nests associated with cysts, one condition should come to mind:

What's the diagnosis?

Melanocytic nevi almost always appear on the bulbar conj during childhood. As with cutaneous nevi, they evolve through a series of histologic conformations, from (in order) junctional to compound to stromal. Conj nevi have low malignant potential.

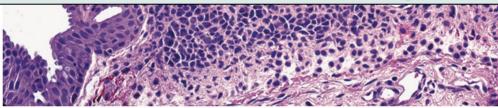




Skin/conj? Conj

There are a couple of field marks that, taken together, nail the diagnosis. What are they?

For more on melanocytic nevi, see slide-set O9

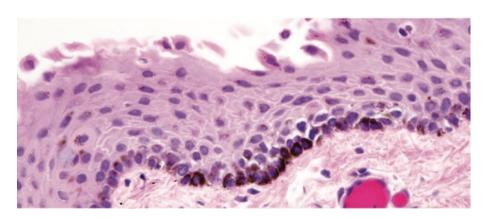


When you encounter melanocytes in nests associated with cysts, one condition should come to mind:

What's the diagnosis?

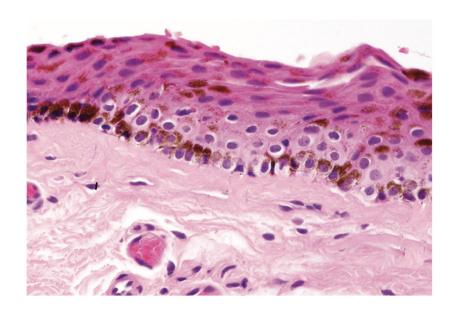
Melanocytic nevi almost always appear on the bulbar conj during childhood. As with cutaneous nevi, they evolve through a series of histologic conformations, from (in order) junctional to compound to stromal. Conj nevi have low malignant potential.



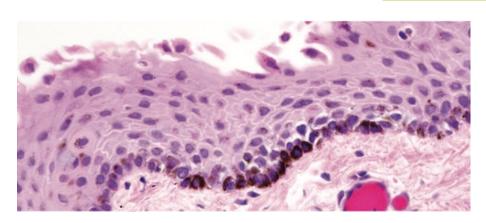


These both represent the same tissue, with relatedbut-different diagnoses.

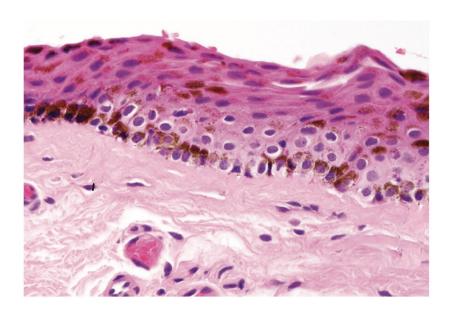
(No question yet—keep going)



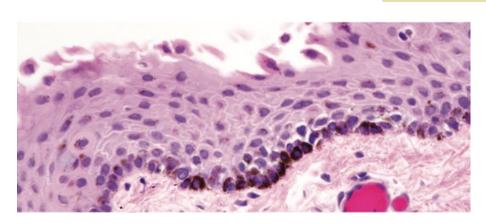




These both represent the same tissue, with related-but-different diagnoses. Skin, or conj?



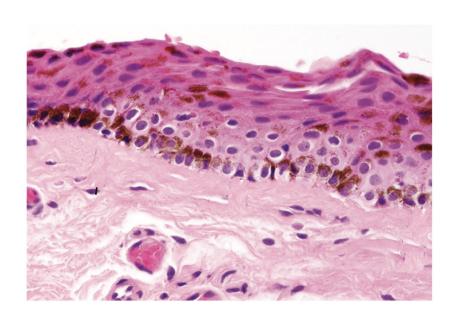




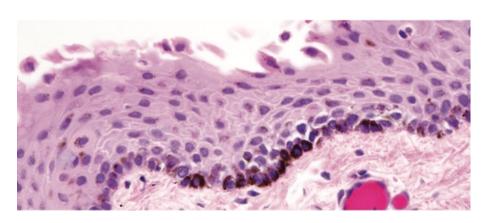
These both represent the same tissue, with relatedbut-different diagnoses.

Skin, or conj?

No keratinization, so conj



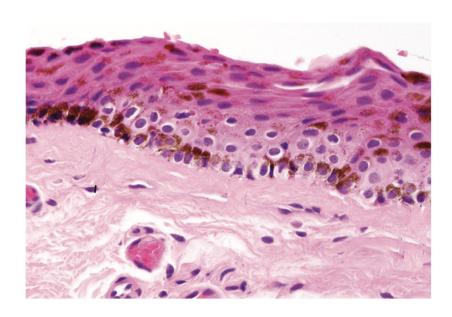




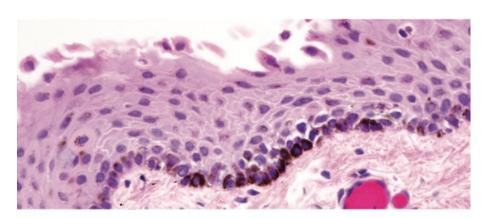
These both represent the same tissue, with relatedbut-different diagnoses.

Skin, or conj?
No keratinization, so conj

What are we supposed to notice in these images?





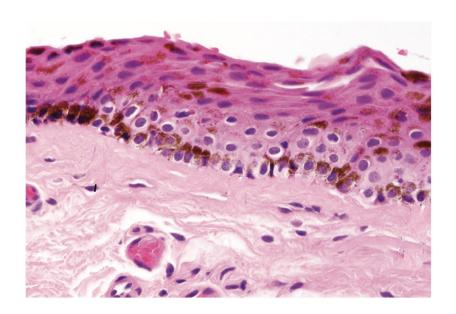


These both represent the same tissue, with relatedbut-different diagnoses.

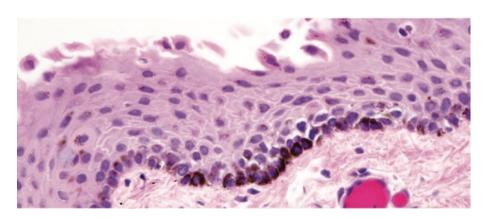
Skin, or conj?
No keratinization, so conj

What are we supposed to notice in these images?

Dat melanin and all those melanocytes. The question being addressed: Can you differentiate between concerning and non-concerning melanocytic lesions?







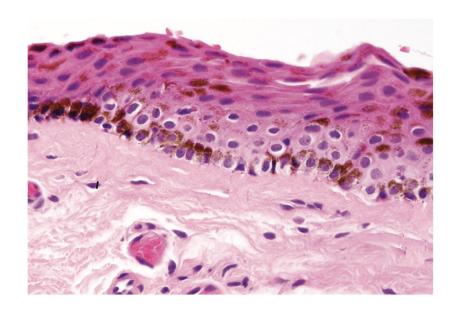
These both represent the same tissue, with relatedbut-different diagnoses.

Skin, or conj?
No keratinization, so conj

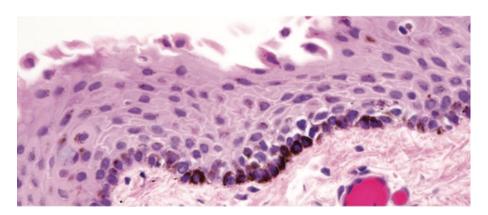
What are we supposed to notice in these images?

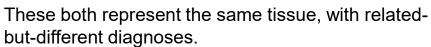
Dat melanin and all those melanocytes. The question being addressed: Can you differentiate between concerning and non-concerning melanocytic lesions?

While oversimplified, we can think of melanocytic conj lesions as falling into one of categories:







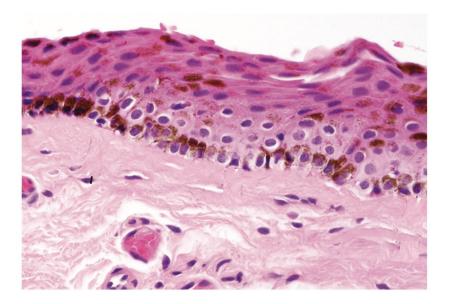


Skin, or conj?
No keratinization, so conj

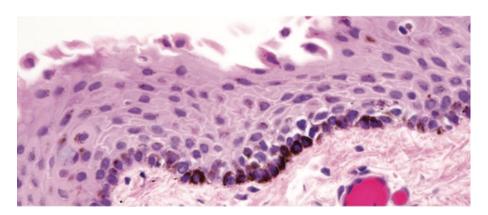
What are we supposed to notice in these images?

Dat melanin and all those melanocytes. The question being addressed: Can you differentiate between concerning and non-concerning melanocytic lesions? While oversimplified, we can think of melanocytic conj lesions as falling into one of five categories:

- --?
- --?
- __2
- --2
- --?







These both represent the same tissue, with relatedbut-different diagnoses.

Skin, or conj?
No keratinization, so conj

What are we supposed to notice in these images?

Dat melanin and all those melanocytes. The question being addressed: Can you differentiate between concerning and non-concerning melanocytic lesions? While oversimplified, we can think of melanocytic conj lesions as falling into one of five categories:

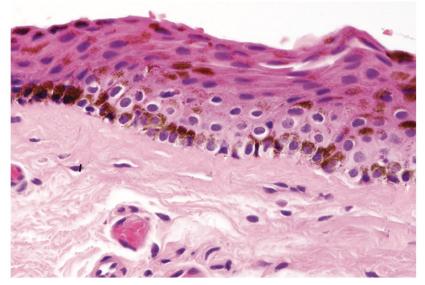
- --Nevus (already addressed)
- --CAM*
- --PAM** without

lotsa words

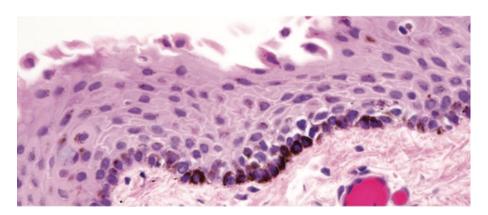
--PAM with

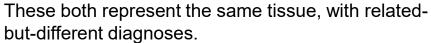
lotsa words

--Melanoma







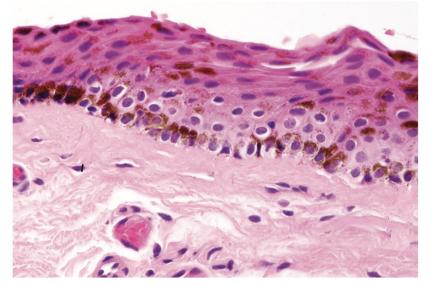


Skin, or conj?
No keratinization, so conj

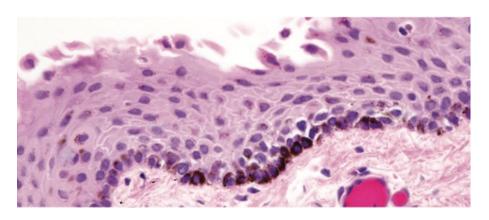
What are we supposed to notice in these images?

Dat melanin and all those melanocytes. The question being addressed: Can you differentiate between concerning and non-concerning melanocytic lesions? While oversimplified, we can think of melanocytic conj lesions as falling into one of five categories:

- --Nevus (already addressed)
- --CAM*
- --PAM** without atypia, or with minimal atypia
- --PAM with moderate to severe atypia
- --Melanoma









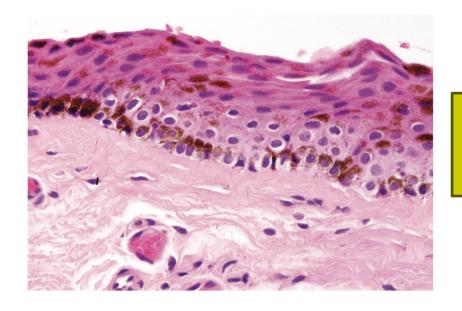
Skin, or conj?

No keratinization, so conj

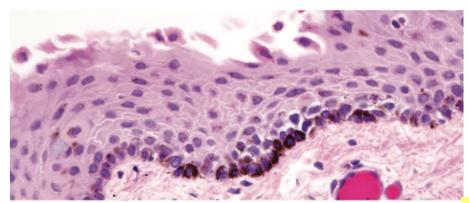
What are we supposed to notice in these images?

Dat melanin and all those melanocytes. The question being addressed: Can you differentiate between concerning and non-concerning melanocytic lesions? While oversimplified, we can think of melanocytic conj lesions as falling into one of five categories:

- --Nevus (already addressed)
- --CAM*
- --PAM** without atypia, or with minimal atypia
- --PAM with moderate to severe atypia







These both represent the same tissue, with relatedbut-different diagnoses.

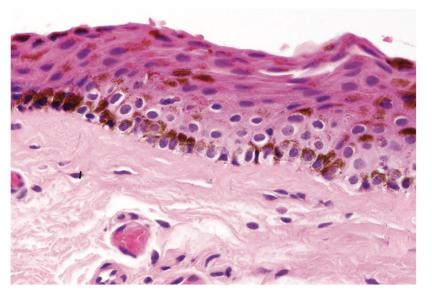
Skin, or conj?

No keratinization, so conj

What are we supposed to notice in these images?

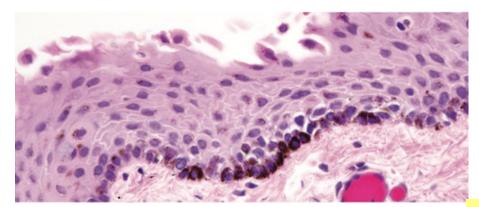
Dat melanin and all those melanocytes. The question

So what are we looking at here? Note that in both images the melanocytes are confined mainly to the basal layer of the epithelium*



--PAIVI WITH moderate to severe atypia





These both represent the same tissue, with relatedbut-different diagnoses.

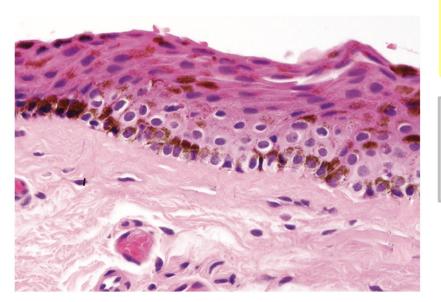
Skin, or conj?

No keratinization, so conj

What are we supposed to notice in these images?

Dat melanin and all those melanocytes. The question

So what are we looking at here? Note that in both images the melanocytes are confined mainly to the basal layer of the epithelium*

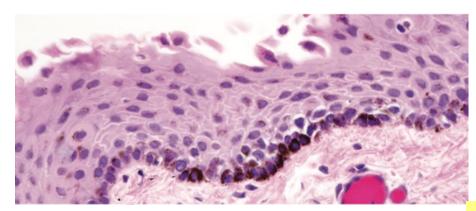


--PAIVI WITH moderate to severe atypia

With regard to malignant potential: CAM and PAM with no/minimal atypia have essentially none, whereas PAM with moderate or severe atypia do.

*Which is not to say that **melanin** is confined to the basal layer—remember, melanocytes package their melanin in melanosomes for distribution to neighboring epi cells





These both represent the same tissue, with relatedbut-different diagnoses.

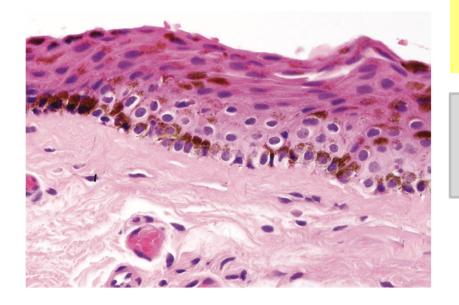
Skin, or conj?

No keratinization, so conj

What are we supposed to notice in these images?

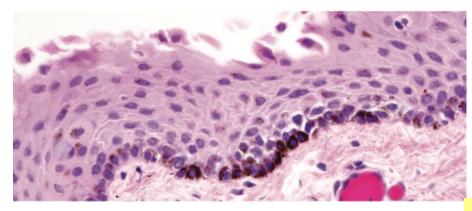
Dat melanin and all those melanocytes. The question

So what are we looking at here? Note that in both images the melanocytes are confined mainly to the basal layer of the epithelium*, and their appearance could be described as typical—or if you'll forgive a double negative, not atypical.*

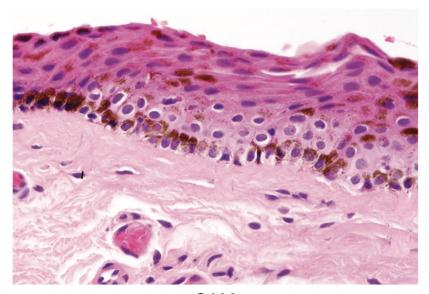


--PAIVI WITH MODERATE TO SEVERE ATYPIA





PAM without atypia



CAM

These both represent the same tissue, with relatedbut-different diagnoses.

Skin, or conj?

No keratinization, so conj

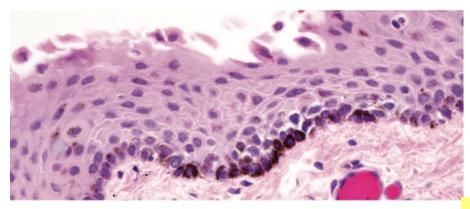
What are we supposed to notice in these images?

Dat melanin and all those melanocytes. The question

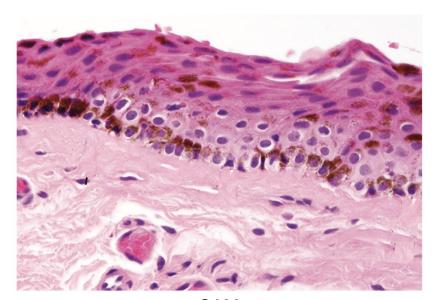
So what are we looking at here? Note that in both images the melanocytes are confined mainly to the basal layer of the epithelium*, and their appearance could be described as typical—or if you'll forgive a double negative, not atypical.* Benign-looking melanocytes confined to the basal epi layer is consistent with both CAM and PAM without/with minimal atypia.

--PAIVI WITH moderate to severe atypia





PAM without atypia



CAM

These both represent the same tissue, with relatedbut-different diagnoses.

Skin, or conj?

No keratinization, so conj

What are we supposed to notice in these images?

Dat melanin and all those melanocytes. The question

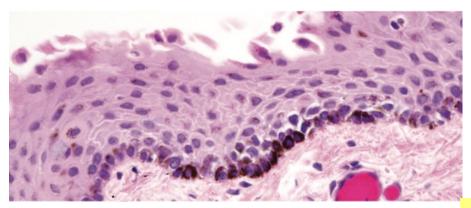
So what are we looking at here? Note that in both

How am I supposed to tell them apart?

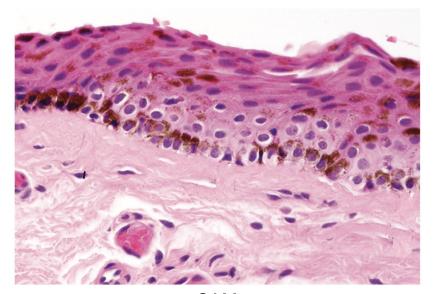
with both CAM and PAM without/with minimal atypia.

--PAIVI WITH moderate to severe atypia





PAM without atypia



CAM

These both represent the same tissue, with relatedbut-different diagnoses.

Skin, or conj?

No keratinization, so conj

What are we supposed to notice in these images?

Dat melanin and all those melanocytes. The question

So what are we looking at here? Note that in both

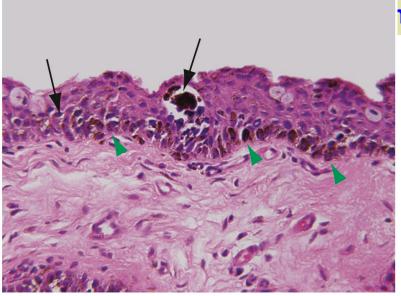
How am I supposed to tell them apart? You're not—not from an H&E slide, anyway. If such a distinction is expected on a test, it would likely be based on clinical info.

with both CAM and PAM without/with minimal atypia -- PAIVI with moderate to severe atypia

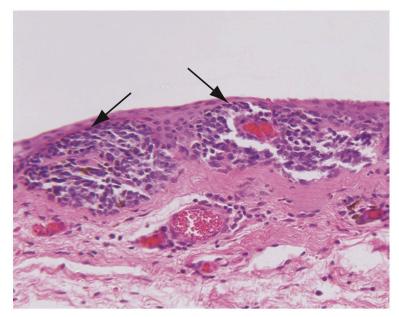




These are what melanocytic badness look like.

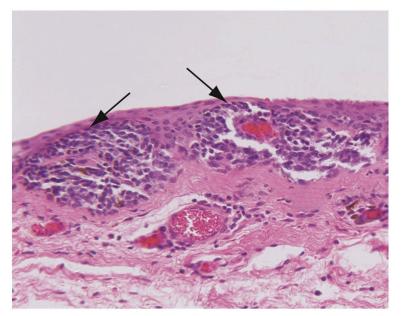


PAM with moderate atypia



PAM with severe atypia (melanoma in situ)

PAM with moderate atypia



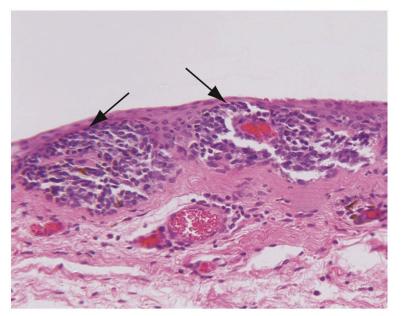
PAM with severe atypia (melanoma in situ)



These are what melanocytic badness look like.

In the *PAM with moderate atypia* pic, note that most of the melanocytes are confined to the basal layer (*arrowheads*). That's a good thing.

PAM with moderate atypia

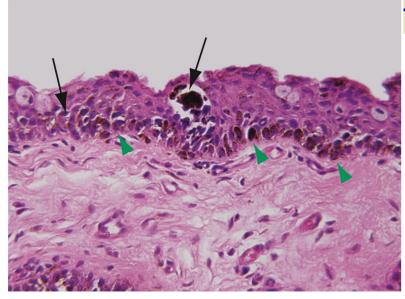


PAM with severe atypia (melanoma in situ)

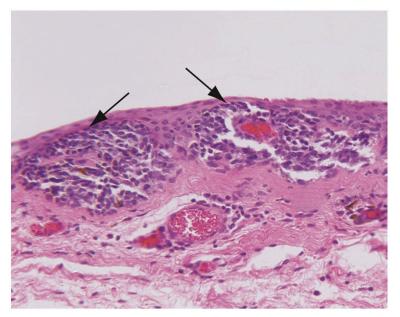


These are what melanocytic badness look like.

In the *PAM with moderate atypia* pic, note that most of the melanocytes are confined to the basal layer (*arrowheads*). That's a good thing. Unfortunately, not all of the melanocytes are basal—some are well up into the more superficial layers (*arrows*). Mos def **not** a good thing.



PAM with moderate atypia



PAM with severe atypia (melanoma in situ)

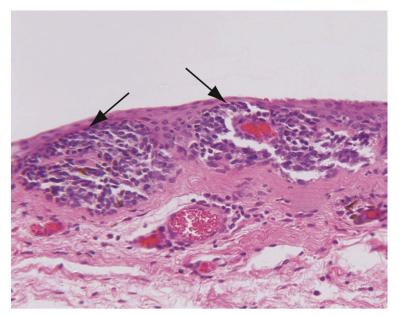


These are what melanocytic badness look like.

In the *PAM with moderate atypia* pic, note that most of the melanocytes are confined to the basal layer (*arrowheads*). That's a good thing. Unfortunately, not all of the melanocytes are basal—some are well up into the more superficial layers (*arrows*). Mos def **not** a good thing.

In the *PAM with severe atypia* pic, the melanocytic proliferation extends into the superficial epithelium (*arrows*)—an ominous finding.

PAM with moderate atypia



PAM with severe atypia (melanoma in situ)



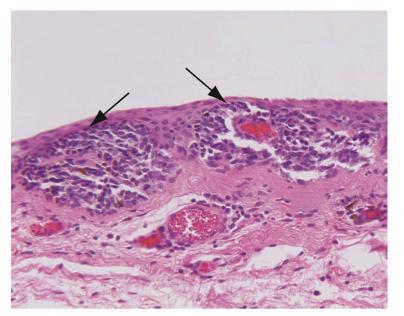
These are what melanocytic badness look like.

In the *PAM with moderate atypia* pic, note that most of the melanocytes are confined to the basal layer (*arrowheads*). That's a good thing. Unfortunately, not all of the melanocytes are basal—some are well up into the more superficial layers (*arrows*). Mos def **not** a good thing.

In the *PAM with severe atypia* pic, the melanocytic proliferation extends into the superficial epithelium (*arrows*)—an ominous finding.

An important aside: Something seems to be missing from the severe atypia pic—what is it?

PAM with moderate atypia



PAM with severe atypia (melanoma in situ)



These are what melanocytic badness look like.

In the *PAM with moderate atypia* pic, note that most of the melanocytes are confined to the basal layer (*arrowheads*). That's a good thing. Unfortunately, not all of the melanocytes are basal—some are well up into the more superficial layers (*arrows*). Mos def **not** a good thing.

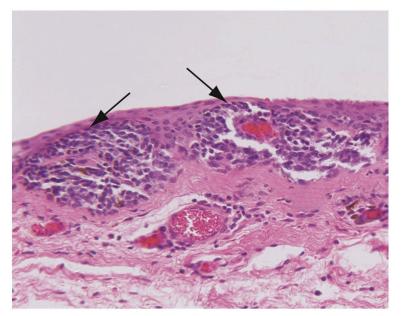
In the *PAM with severe atypia* pic, the melanocytic proliferation extends into the superficial epithelium (*arrows*)—an ominous finding.

An important aside: Something seems to be missing from the severe atypia pic—what is it?

Melanin—there's hardly any present at all. Clinically, this lesion was

.

PAM with moderate atypia



PAM with severe atypia (melanoma in situ)



These are what melanocytic badness look like.

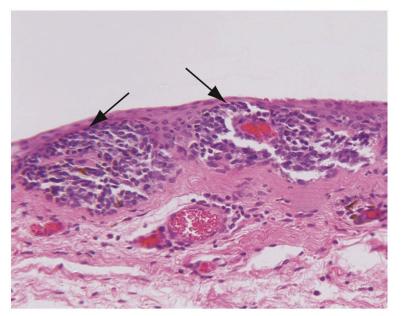
In the *PAM with moderate atypia* pic, note that most of the melanocytes are confined to the basal layer (*arrowheads*). That's a good thing. Unfortunately, not all of the melanocytes are basal—some are well up into the more superficial layers (*arrows*). Mos def **not** a good thing.

In the *PAM with severe atypia* pic, the melanocytic proliferation extends into the superficial epithelium (*arrows*)—an ominous finding.

An important aside: Something seems to be missing from the severe atypia pic—what is it?

Melanin—there's hardly any present at all. Clinically, this lesion was amelanotic.

PAM with moderate atypia



PAM with severe atypia (melanoma in situ)



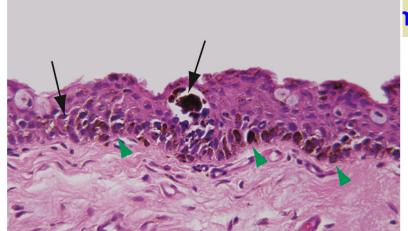
These are what melanocytic badness look like.

In the *PAM with moderate atypia* pic, note that most of the melanocytes are confined to the basal layer (*arrowheads*). That's a good thing. Unfortunately, not all of the melanocytes are basal—some are well up into the more superficial layers (*arrows*). Mos def **not** a good thing.

In the *PAM with severe atypia* pic, the melanocytic proliferation extends into the superficial epithelium (*arrows*)—an ominous finding.

An important aside: Something seems to be missing from the severe atypia pic—what is it?

Melanin—there's hardly any present at all. Clinically, this lesion was amelanotic. The point being, don't depend on the presence of melanin to pull the trigger on a melanocytic diagnosis!

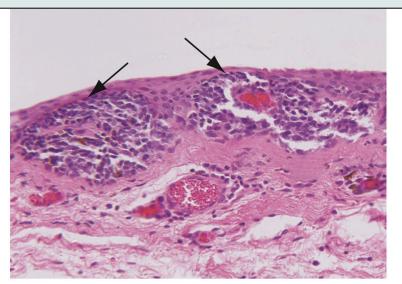




These are what melanocytic badness look like.

In the *PAM with moderate atypia* pic, note that most of the melanocytes are confined to the basal layer (*arrowheads*). That's a good thing. Unfortunately, not all of the melanocytes are basal—some are well up into the more superficial layers (*arrows*). Mos def **not** a good thing.

For more on CAM and PAM, see slide-set O9



PAM with severe atypia (melanoma in situ)

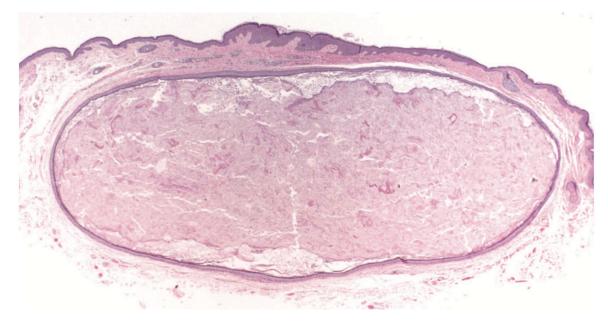
(arrows)—an ominous finding.

An important aside: Something seems to be missing from the severe atypia pic—what is it?

Melanin—there's hardly any present at all. Clinically, this lesion was amelanotic. The point being, don't depend on the presence of melanin to pull the trigger on a melanocytic diagnosis!



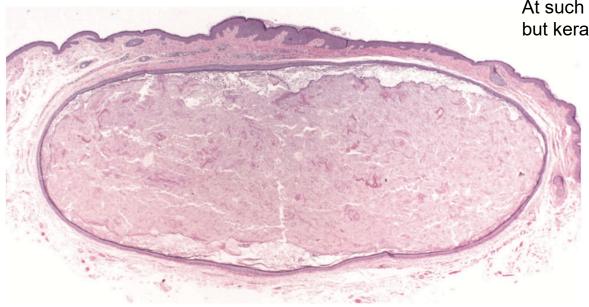
Skin, or conj?





Skin, or conj?

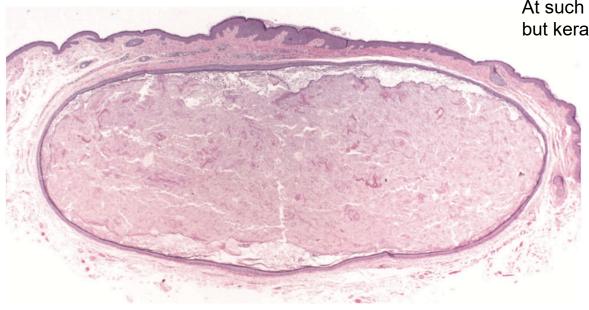
At such a low power it's tough to see, but keratinization is present





Skin, or conj?

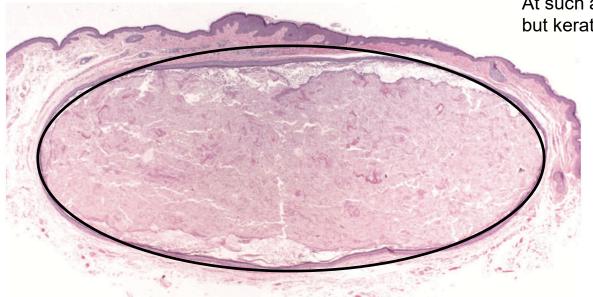
At such a low power it's tough to see, but keratinization is present, so skin





Skin, or conj?

At such a low power it's tough to see, but keratinization is present, so skin

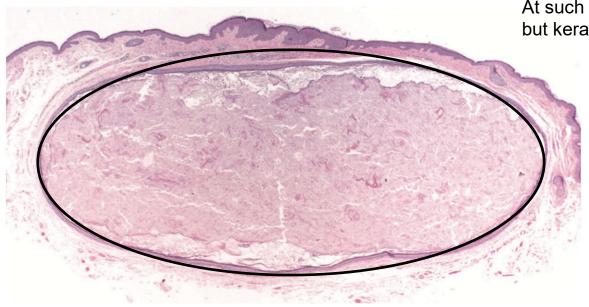


So what's going on here? The most obvious finding is **this** large something-like structure.



Skin, or conj?

At such a low power it's tough to see, but keratinization is present, so skin

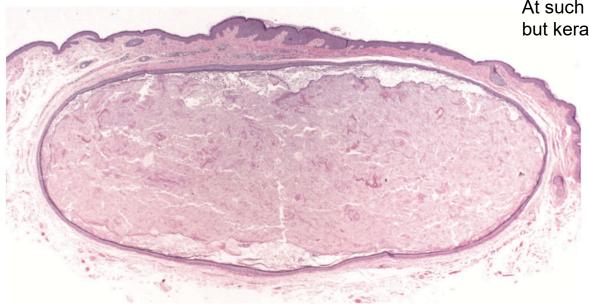


So what's going on here? The most obvious finding is **this** large cyst-like structure.



Skin, or conj?

At such a low power it's tough to see, but keratinization is present, so skin

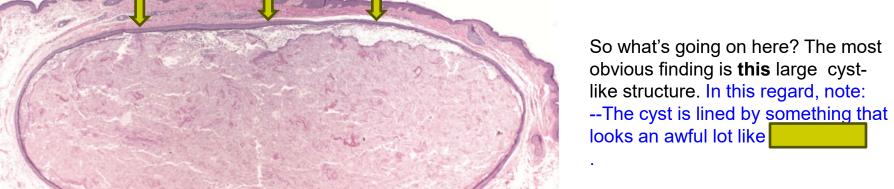


So what's going on here? The most obvious finding is **this** large cyst-like structure. In this regard, note:



Skin, or conj?

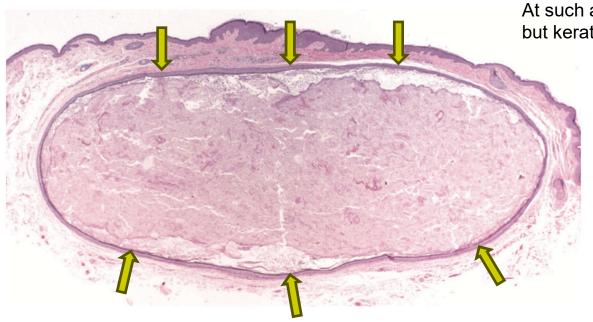
At such a low power it's tough to see, but keratinization is present, so skin





Skin, or conj?

At such a low power it's tough to see, but keratinization is present, so skin

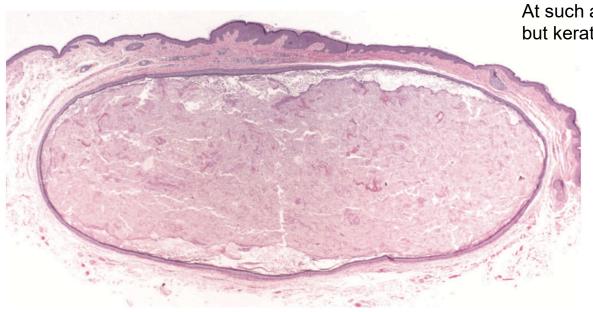


So what's going on here? The most obvious finding is **this** large cyst-like structure. In this regard, note:
--The cyst is lined by something that looks an awful lot like epithelium.



Skin, or conj?

At such a low power it's tough to see, but keratinization is present, so skin

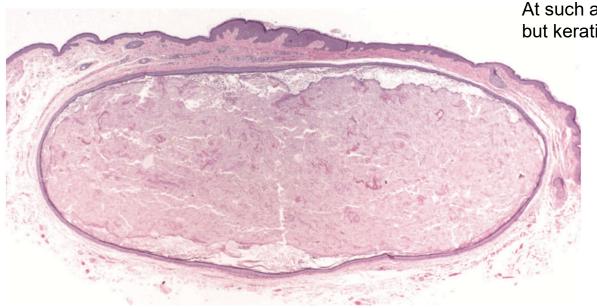


So what's going on here? The most obvious finding is **this** large cyst-like structure. In this regard, note:
--The cyst is lined by something that looks an awful lot like epithelium.
--The cyst contains a fairly uniform, amorphous and acellular material.



Skin, or conj?

At such a low power it's tough to see, but keratinization is present, so skin



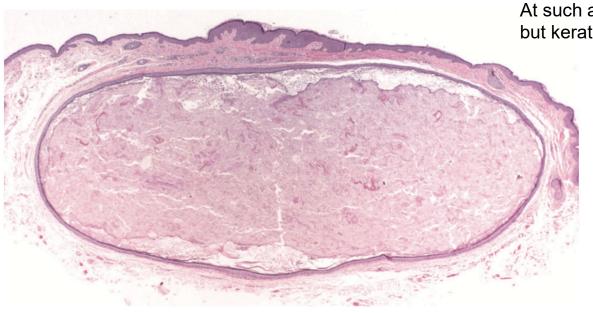
So what's going on here? The most obvious finding is **this** large cyst-like structure. In this regard, note:
--The cyst is lined by something that looks an awful lot like epithelium.

- --The cyst contains a fairly uniform, amorphous and acellular material.
- --The cyst is located in the



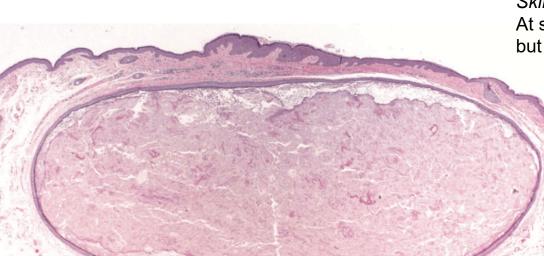
Skin, or conj?

At such a low power it's tough to see, but keratinization is present, so skin



So what's going on here? The most obvious finding is **this** large cyst-like structure. In this regard, note:
--The cyst is lined by something that looks an awful lot like epithelium.
--The cyst contains a fairly uniform, amorphous and acellular material.
--The cyst is located in the dermis.





Skin, or conj?
At such a low power it's tough to see,

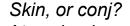
but keratinization is present, so skin

So what's going on here? The most obvious finding is **this** large cyst-like structure. In this regard, note:
--The cyst is lined by something that looks an awful lot like epithelium.
--The cyst contains a fairly uniform, amorphous and acellular material.
--The cyst is located in the dermis.

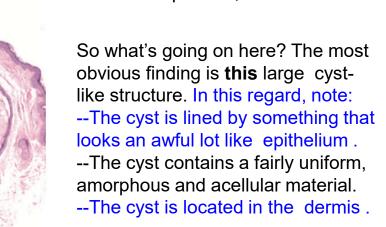
What's the diagnosis?

When you encounter a cyst in the dermis lined with epi and containing an amorphous material, one dx should come to mind:





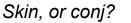
At such a low power it's tough to see, but keratinization is present, so skin



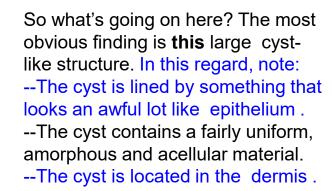
When you encounter a cyst in the dermis lined with epi and containing an amorphous material, one dx should come to mind:

Epidermal inclusion cyst (aka cyst) is a common lid finding.





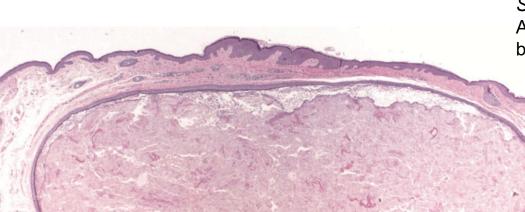
At such a low power it's tough to see, but keratinization is present, so skin



When you encounter a cyst in the dermis lined with epi and containing an amorphous material, one dx should come to mind:

Epidermal inclusion cyst (aka epidermoid cyst) is a common lid finding.





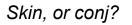
Skin, or conj?
At such a low power it's tough to see,
but keratinization is present, so skin

So what's going on here? The most obvious finding is **this** large cyst-like structure. In this regard, note:
--The cyst is lined by something that looks an awful lot like epithelium.
--The cyst contains a fairly uniform, amorphous and acellular material.
--The cyst is located in the dermis.

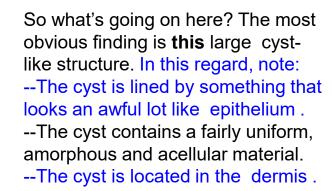
When you encounter a cyst in the dermis lined with epi and containing an amorphous material, one dx should come to mind:

Epidermal inclusion cyst (aka *epidermoid* cyst) is a common lid finding. Like the lid itself, the cyst is lined with layering shape action epithelium.





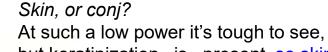
At such a low power it's tough to see, but keratinization is present, so skin



When you encounter a cyst in the dermis lined with epi and containing an amorphous material, one dx should come to mind:

Epidermal inclusion cyst (aka *epidermoid* cyst) is a common lid finding. Like the lid itself, the cyst is lined with stratified squamous keratinizing epithelium.



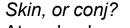


but keratinization is present, so skin

So what's going on here? The most obvious finding is **this** large cyst-like structure. In this regard, note:
--The cyst is lined by something that looks an awful lot like epithelium.
--The cyst contains a fairly uniform, amorphous and acellular material.
--The cyst is located in the dermis.

When you encounter a cyst in the dermis lined with epi and containing an amorphous material, one dx should come to mind:





At such a low power it's tough to see, but keratinization is present, so skin

So what's going on here? The most obvious finding is **this** large cyst-like structure. In this regard, note:
--The cyst is lined by something that looks an awful lot like epithelium.
--The cyst contains a fairly uniform, amorphous and acellular material.
--The cyst is located in the dermis.

When you encounter a cyst in the dermis lined with epi and containing an amorphous material, one dx should come to mind:



nat

Skin, or conj?
At such a low power it's tough to see,
but keratinization is present, so skin

Since it's in the dermis, shouldn't it be a dermal cyst?

When you encounter a cyst in the dermis lined with epi and containing an amorphous material, one dx should come to mind:



nat

Skin, or conj?
At such a low power it's tough to see,
but keratinization is present, so skin

Since it's in the dermis, shouldn't it be a dermal cyst?
You'd think so, but no. While similar to epidermal cysts, dermoid cysts differ in that they contain two words

When you encounter a cyst in the dermis lined with epi and containing an amorphous material, one dx should come to mind:



nat

Skin, or conj?
At such a low power it's tough to see,
but keratinization is present, so skin

Since it's in the dermis, shouldn't it be a **dermal** cyst? You'd think so, but no. While similar to epidermal cysts, dermoid cysts differ in that they contain dermal appendages

When you encounter a cyst in the dermis lined with epi and containing an amerphous material, one dx should come to mind:



nat

Skin, or conj?
At such a low power it's tough to see,
but keratinization is present, so skin

Since it's in the dermis, shouldn't it be a **dermal** cyst?

You'd think so, but no. While similar to epidermal cysts, dermoid cysts differ in that they contain dermal appendages (aka two words

When you encounter a cyst in the dermis lined with epi and containing an amorphous

material, one dx should come to mind:



nat

Skin, or conj?
At such a low power it's tough to see,
but keratinization is present, so skin

Since it's in the dermis, shouldn't it be a **dermal** cyst? You'd think so, but no. While similar to epidermal cysts, dermoid cysts differ in that they contain dermal appendages (aka adnexal structures)

When you encounter a **cyst in the dermis** lined with epi and containing an amorphous

material, one dx should come to mind:



st

nat

Skin, or conj?
At such a low power it's tough to see,
but keratinization is present, so skin

Since it's in the dermis, shouldn't it be a **dermal** cyst?

You'd think so, but no. While similar to epidermal cysts, dermoid cysts differ in that they contain dermal appendages (aka adnexal structures) such as two words and two words.

When you encounter a cyst in the dermis lined with epi and containing an amorphous material, one dx should come to mind:



Skin, or conj?
At such a low power it's tough to see,
but keratinization is present, so skin

Since it's in the dermis, shouldn't it be a dermal cyst?

You'd think so, but no. While similar to epidermal cysts, dermoid cysts differ in that they contain dermal appendages (aka adnexal structures) such as hair follicles and sebaceous glands.

When you encounter a cyst in the dermis lined with epi and containing an amerphous material, one dx should come to mind:

Epidermal inclusion cyst (aka *epidermoid* cyst) is a common lid finding. Like the lid itself, the cyst is lined with stratified squamous keratinizing epithelium. (This explains why the amorphous material filling the cyst is keratin.)

nat

· .



nat

Skin, or conj?
At such a low power it's tough to see,
but keratinization is present, so skin

Since it's in the dermis, shouldn't it be a **dermal** cyst?

You'd think so, but no. While similar to epidermal cysts, dermoid cysts differ in that they contain dermal appendages (aka adnexal structures) such as hair follicles and sebaceous glands. Thus, the lumen of a dermoid cyst will contain and

When you encounter a cyst in the dermis lined with epi and containing an amerphous material, one dx should come to mind:



Skin, or conj?
At such a low power it's tough to see,
but keratinization is present, so skin

Since it's in the dermis, shouldn't it be a dermal cyst?

You'd think so, but no. While similar to epidermal cysts, dermoid cysts differ in that they contain dermal appendages (aka adnexal structures) such as hair follicles and sebaceous glands. Thus, the lumen of a dermoid cyst will contain hair and sebum

When you encounter a cyst in the dermis lined with epi and containing an amerphous material, one dx should come to mind:

Epidermal inclusion cyst (aka *epidermoid* cyst) is a common lid finding. Like the lid itself, the cyst is lined with stratified squamous keratinizing epithelium. (This explains why the amorphous material filling the cyst is keratin.)

nat

.



nat

Skin, or conj?
At such a low power it's tough to see,
but keratinization is present, so skin

Since it's in the dermis, shouldn't it be a dermal cyst?

You'd think so, but no. While similar to epidermal cysts, dermoid cysts differ in that they contain dermal appendages (aka adnexal structures) such as hair follicles and sebaceous glands. Thus, the lumen of a dermoid cyst will contain hair and sebum (in addition to _____).

When you encounter a cyst in the dermis lined with epi and containing an amerphous material, one dx should come to mind:



Skin, or conj?
At such a low power it's tough to see,
but keratinization is present, so skin

Since it's in the dermis, shouldn't it be a dermal cyst?

You'd think so, but no. While similar to epidermal cysts, dermoid cysts differ in that they contain dermal appendages (aka adnexal structures) such as hair follicles and sebaceous glands. Thus, the lumen of a dermoid cyst will contain hair and sebum (in addition to keratin).

When you encounter a cyst in the dermis lined with epi and containing an amorphous material, one dx should come to mind:

Epidermal inclusion cyst (aka *epidermoid* cyst) is a common lid finding. Like the lid itself, the cyst is lined with stratified squamous keratinizing epithelium. (This explains why the amorphous material filling the cyst is keratin.)

nat



nat

Skin, or conj?
At such a low power it's tough to see,
but keratinization is present, so skin

Since it's in the dermis, shouldn't it be a dermal cyst?

You'd think so, but no. While similar to epidermal cysts, dermoid cysts differ in that they contain dermal appendages (aka *adnexal structures*) such as hair follicles and sebaceous glands. Thus, the lumen of a dermoid cyst will contain hair and sebum (in addition to keratin). It is the presence of dermal appendages that give the dermoid cyst its name.

When you encounter a cyst in the dermis lined with epi and containing an amerphous material, one dx should come to mind:



Skin, or conj?
At such a low power it's tough to see,
but keratinization is present, so skin

Since it's in the dermis, shouldn't it be a dermal cyst?

You'd think so, but no. While similar to epidermal cysts, dermoid cysts differ in that they contain dermal appendages (aka *adnexal structures*) such as hair follicles and sebaceous glands. Thus, the lumen of a dermoid cyst will contain hair and sebum (in addition to keratin). It is the presence of dermal appendages that give the dermoid cyst its name.

Are we going to see a photomicrograph of a dermoid cyst?

When you encounter a cyst in the dermis lined with epi and containing an amerphous material, one dx should come to mind:

Epidermal inclusion cyst (aka *epidermoid* cyst) is a common lid finding. Like the lid itself, the cyst is lined with stratified squamous keratinizing epithelium. (This explains why the amorphous material filling the cyst is keratin.)

nat

١,

.



nat

Skin, or conj?
At such a low power it's tough to see,
but keratinization is present, so skin

Since it's in the dermis, shouldn't it be a dermal cyst?

You'd think so, but no. While similar to epidermal cysts, dermoid cysts differ in that they contain dermal appendages (aka *adnexal structures*) such as hair follicles and sebaceous glands. Thus, the lumen of a dermoid cyst will contain hair and sebum (in addition to keratin). It is the presence of dermal appendages that give the dermoid cyst its name.

Are we going to see a photomicrograph of a dermoid cyst?

Not in this section. While they can occur in the lids, they are much more common in the

When you encounter a cyst in the dermis lined with epi and containing an amorphous material, one dx should come to mind:



nat

Skin, or conj?
At such a low power it's tough to see,
but keratinization is present, so skin

Since it's in the dermis, shouldn't it be a dermal cyst?

You'd think so, but no. While similar to epidermal cysts, dermoid cysts differ in that they contain dermal appendages (aka *adnexal structures*) such as hair follicles and sebaceous glands. Thus, the lumen of a dermoid cyst will contain hair and sebum (in addition to keratin). It is the presence of dermal appendages that give the dermoid cyst its name.

Are we going to see a photomicrograph of a dermoid cyst?

Not in this section. While they can occur in the lids, they are much more common in the orbit.

When you encounter a cyst in the dermis lined with epi and containing an amorphous material, one dx should come to mind:

Epidermal inclusion cyst (aka *epidermoid* cyst) is a common lid finding. Like the lid itself, the cyst is lined with stratified squamous keratinizing epithelium. (This explains why the amorphous material filling the cyst is keratin.)



nat

Skin, or conj?
At such a low power it's tough to see,
but keratinization is present, so skin

Since it's in the dermis, shouldn't it be a dermal cyst?

You'd think so, but no. While similar to epidermal cysts, dermoid cysts differ in that they contain dermal appendages (aka *adnexal structures*) such as hair follicles and sebaceous glands. Thus, the lumen of a dermoid cyst will contain hair and sebum (in addition to keratin). It is the presence of dermal appendages that give the dermoid cyst its name.

Are we going to see a photomicrograph of a dermoid cyst?

Not in this section. While they can occur in the lids, they are much more common in the orbit. (Classic presentation: A finding in the quadrant orbit of a stage)

When you encounter a cyst in the dermis lined with epi and containing an amerphous material, one dx should come to mind:

Epidermal inclusion cyst (aka *epidermoid* cyst) is a common lid finding. Like the lid itself, the cyst is lined with stratified squamous keratinizing epithelium. (This explains why the amorphous material filling the cyst is keratin.)



nat

Skin, or conj?
At such a low power it's tough to see,
but keratinization is present, so skin

Since it's in the dermis, shouldn't it be a dermal cyst?

You'd think so, but no. While similar to epidermal cysts, dermoid cysts differ in that they contain dermal appendages (aka *adnexal structures*) such as hair follicles and sebaceous glands. Thus, the lumen of a dermoid cyst will contain hair and sebum (in addition to keratin). It is the presence of dermal appendages that give the dermoid cyst its name.

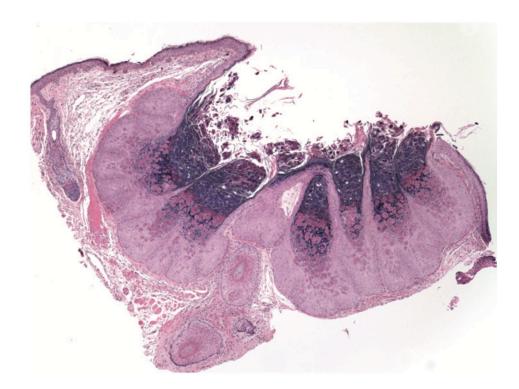
Are we going to see a photomicrograph of a dermoid cyst?

Not in this section. While they can occur in the lids, they are much more common in the orbit. (Classic presentation: A mass in the superotemporal orbit of a child.)

When you encounter a cyst in the dermis lined with epi and containing an amerphous material, one dx should come to mind:

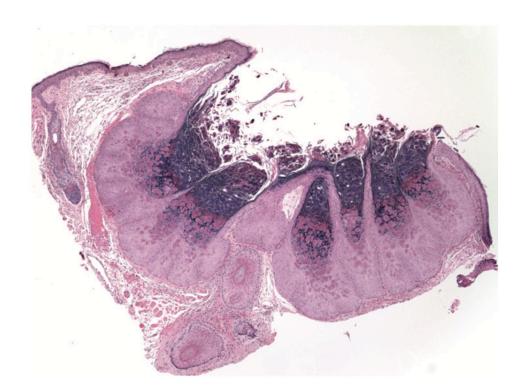
Epidermal inclusion cyst (aka *epidermoid* cyst) is a common lid finding. Like the lid itself, the cyst is lined with stratified squamous keratinizing epithelium. (This explains why the amorphous material filling the cyst is keratin.)





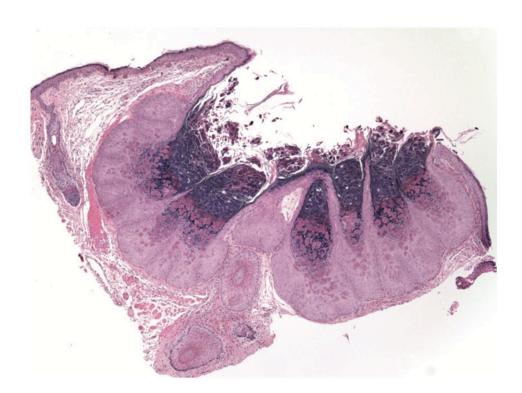
Skin, or conj?





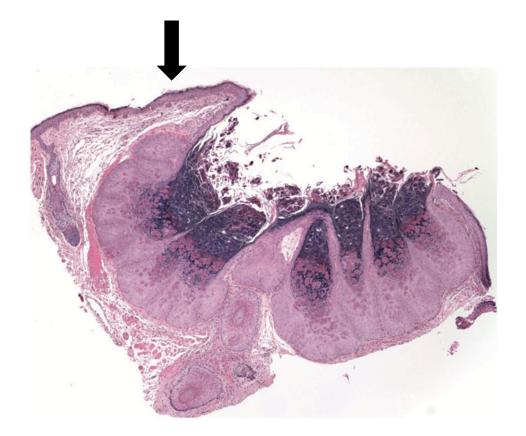
Skin, or conj?
Again, low mag makes it a tough call.





Skin, or conj?
Again, low mag makes it a tough call. That said, keratinization is y present





Skin, or conj?
Again, low mag makes it a tough call. That said, keratinization is present, so skin.





Skin, or conj?
Again, low mag makes it a tough call. That said, keratinization is present, so skin.

So what are we to make of this? First, note the unusual shape of the lesion.





Skin, or conj?
Again, low mag makes it a tough call. That said, keratinization is present, so skin.

So what are we to make of this? First, note the unusual shape of the lesion. (The term of art for its shape is two-words .)

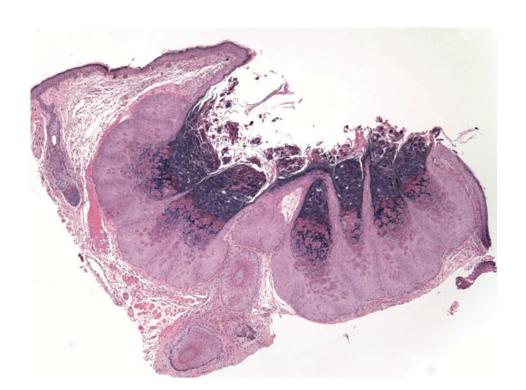




Skin, or conj?
Again, low mag makes it a tough call. That said, keratinization is present, so skin.

So what are we to make of this? First, note the unusual shape of the lesion. (The term of art for its shape is 'cup-like'.)

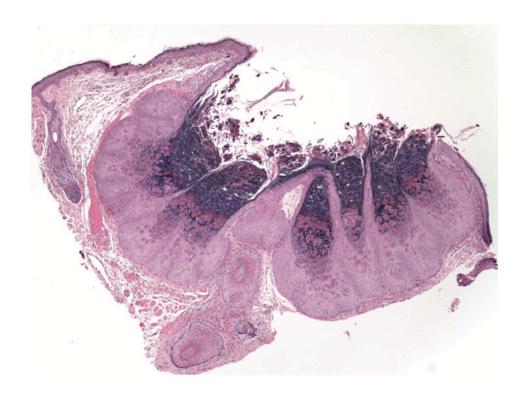




Skin, or conj?
Again, low mag makes it a tough call. That said, keratinization is present, so skin.

So what are we to make of this? First, note the unusual shape of the lesion. (The term of art for its shape is 'cup-like'.) The epithelium at the bottom of the cup is dramatically thickened, and the cup is filled with an amorphous acellular material that seems friable.



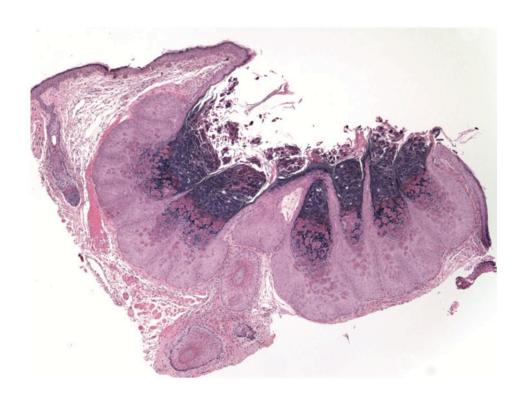


What's the diagnosis?

Skin, or conj?
Again, low mag makes it a tough call. That said, keratinization is present, so skin.

So what are we to make of this? First, note the unusual shape of the lesion. (The term of art for its shape is 'cup-like'.) The epithelium at the bottom of the cup is dramatically thickened, and the cup is filled with an amorphous acellular material that seems friable. A lid lesion with a gradeau-filled 'cup' should bring to mind one lesion in particular:





Skin, or conj?

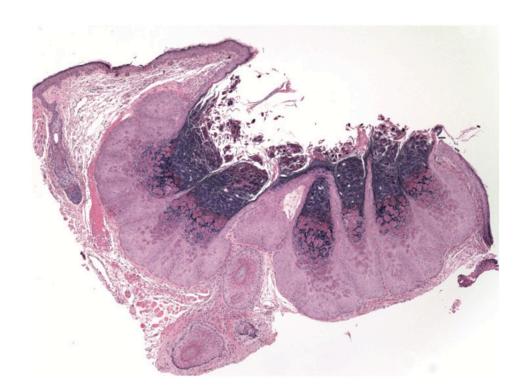
Again, low mag makes it a tough call. That said, keratinization is present, so skin.

So what are we to make of this? First, note the unusual shape of the lesion. (The term of art for its shape is 'cup-like'.) The epithelium at the bottom of the cup is dramatically thickened, and the cup is filled with an amorphous acellular material that seems friable. A lid lesion with a gradeau-filled 'cup' should bring to mind one lesion in particular:

What's the diagnosis?

Molluscum contagiosum lesions arise from a type of infection.





Skin, or conj?

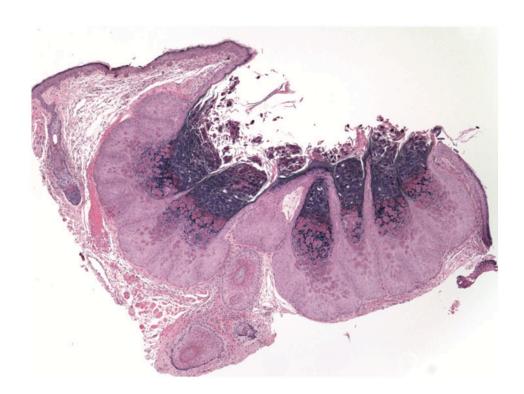
Again, low mag makes it a tough call. That said, keratinization is present, so skin.

So what are we to make of this? First, note the unusual shape of the lesion. (The term of art for its shape is 'cup-like'.) The epithelium at the bottom of the cup is dramatically thickened, and the cup is filled with an amorphous acellular material that seems friable. A lid lesion with a gradeau-filled 'cup' should bring to mind one lesion in particular:

What's the diagnosis?

Molluscum contagiosum lesions arise from a viral infection.





Skin, or conj?

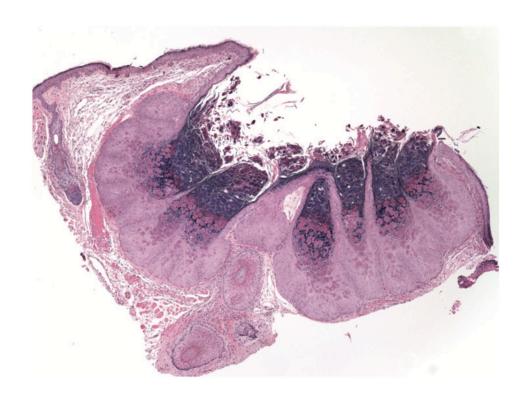
Again, low mag makes it a tough call. That said, keratinization is present, so skin.

So what are we to make of this? First, note the unusual shape of the lesion. (The term of art for its shape is 'cup-like'.) The epithelium at the bottom of the cup is dramatically thickened, and the cup is filled with an amorphous acellular material that seems friable. A lid lesion with a gradeau-filled 'cup' should bring to mind one lesion in particular:

What's the diagnosis?

Molluscum contagiosum lesions arise from a viral infection. Clinically they are shaped nodules





Skin, or conj?
Again, low mag makes it a t

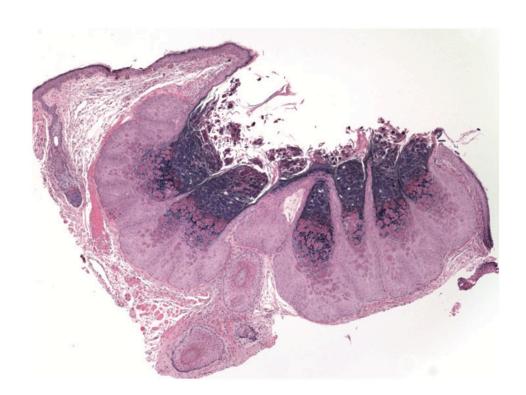
Again, low mag makes it a tough call. That said, keratinization is present, so skin.

So what are we to make of this? First, note the unusual shape of the lesion. (The term of art for its shape is 'cup-like'.) The epithelium at the bottom of the cup is dramatically thickened, and the cup is filled with an amorphous acellular material that seems friable. A lid lesion with a gradeau-filled 'cup' should bring to mind one lesion in particular:

What's the diagnosis?

Molluscum contagiosum lesions arise from a viral infection. Clinically they are dome –shaped nodules





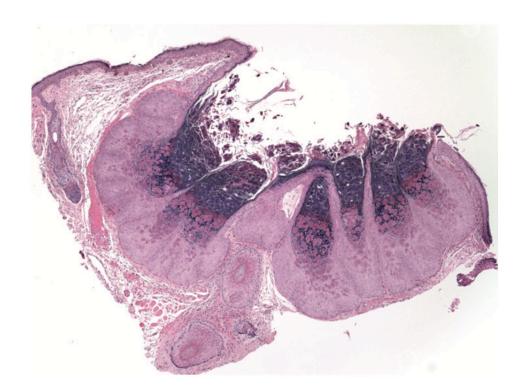
Skin, or conj?
Again, low mag makes it a tough call. That said, keratinization is present, so skin.

So what are we to make of this? First, note the unusual shape of the lesion. (The term of art for its shape is 'cup-like'.) The epithelium at the bottom of the cup is dramatically thickened, and the cup is filled with an amorphous acellular material that seems friable. A lid lesion with a gradeau-filled 'cup' should bring to mind one lesion in particular:

What's the diagnosis?

Molluscum contagiosum lesions arise from a viral infection. Clinically they are dome –shaped nodules with a central term of art (the 'cup' component of the lesion).





What's the diagnosis?

Skin, or conj?
Again, low mag makes it a tough call. That said, keratinization is present, so skin.

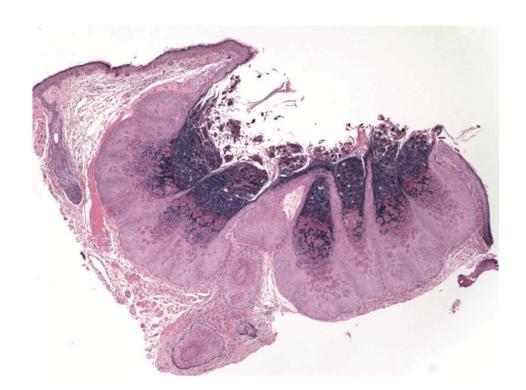
So what are we to make of this? First, note the unusual shape of the lesion. (The term of art for its shape is 'cup-like'.) The epithelium at the bottom of the cup is dramatically thickened, and the cup is filled with an amorphous acellular

material t gradeaulesion in



Molluscum contagiosum lesions arise from a viral infection. Clinically they are dome —shaped nodules with a central umbilication (the 'cup' component of the lesion).





Skin, or conj?
Again, low mag makes it a tough call. That said, keratinization is present, so skin.

So what are we to make of this? First, note the unusual shape of the lesion. (The term of art for its shape is 'cup-like'.) The epithelium at the bottom of the cup is dramatically thickened, and the cup is filled with an amorphous acellular material that seems friable. A lid lesion with a gradeau-filled 'cup' should bring to mind one lesion in particular:

What's the diagnosis?

Molluscum contagiosum lesions arise from a viral infection. Clinically they are dome –shaped nodules with a central umbilication (the 'cup' component of the lesion). Necrotic cells are shed, filling the cup with amorphous funk.



Skin or conj?







*Skin or conj?*Skin





Skin or conj? Skin

What are we supposed to notice? A few things:

- --?
- -- ?
- --?





Skin or conj? Skin

--?

-- ?





Skin or conj? Skin

What are we supposed to notice? A few things: --The overall configuration is dome --shaped

- -- ?
- --?





Skin or conj? Skin

What are we supposed to notice? A few things:

- --The overall configuration is dome -shaped

--?





Skin or conj? Skin

What are we supposed to notice? A few things:

- --The overall configuration is dome -shaped
- --The epi is hyperplastic, and looks benign

--?





Skin or conj? Skin

What are we supposed to notice? A few things:

- --The overall configuration is dome -shaped
- --The epi is hyperplastic, and looks benign
- --The lesion has a number of cysts containing

 distribution (two words)

 collections of surface



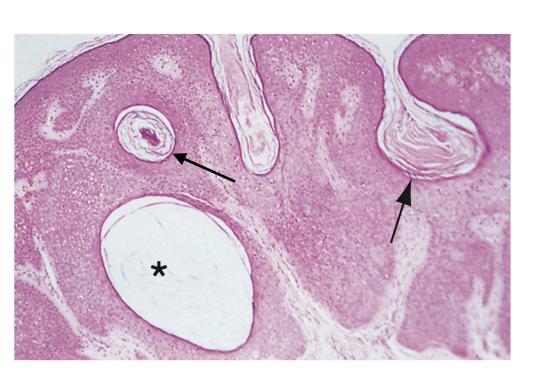


Skin or conj? Skin

What are we supposed to notice? A few things:

- --The overall configuration is dome -shaped
- --The epi is hyperplastic, and looks benign
- --The lesion has a number of cysts containing concentrically laminated collections of surface keratin





Skin or conj? Skin

What are we supposed to notice? A few things:

- --The overall configuration is dome -shaped
- --The epi is hyperplastic, and looks benign
- --The lesion has a number of cysts containing concentrically laminated collections of surface keratin

Here is a close-up wherein the concentric lamination is easier to appreciate (both *asterisk* and *arrows*)





What's the diagnosis?

Skin or conj? Skin

What are we supposed to notice? A few things:

- --The overall configuration is dome -shaped
- --The epi is hyperplastic, and looks benign
- --The lesion has a number of cysts containing concentrically laminated collections of surface keratin

Here is a close-up wherein the concentric lamination is easier to appreciate (both *asterisk* and *arrows*)

If you encounter a dome-shaped lesion containing such cysts, one diagnosis should come to mind:





What's the diagnosis?

Skin or conj? Skin

What are we supposed to notice? A few things:

- --The overall configuration is dome -shaped
- --The epi is hyperplastic, and looks benign
- --The lesion has a number of cysts containing concentrically laminated collections of surface keratin

Here is a close-up wherein the concentric lamination is easier to appreciate (both *asterisk* and *arrows*)

If you encounter a dome-shaped lesion containing such cysts, one diagnosis should come to mind:

Seborrheic keratosis is a



epithelial proliferation





What's the diagnosis?

Skin or conj? Skin

What are we supposed to notice? A few things:

- --The overall configuration is dome -shaped
- --The epi is hyperplastic, and looks benign
- --The lesion has a number of cysts containing concentrically laminated collections of surface keratin

Here is a close-up wherein the concentric lamination is easier to appreciate (both *asterisk* and *arrows*)

If you encounter a dome-shaped lesion containing such cysts, one diagnosis should come to mind:

Seborrheic keratosis is a common epithelial proliferation





What's the diagnosis?

Skin or conj? Skin

What are we supposed to notice? A few things:

- --The overall configuration is dome -shaped
- --The epi is hyperplastic, and looks benign
- --The lesion has a number of cysts containing concentrically laminated collections of surface keratin

Here is a close-up wherein the concentric lamination is easier to appreciate (both *asterisk* and *arrows*)

If you encounter a dome-shaped lesion containing such cysts, one diagnosis should come to mind:

Seborrheic keratosis is a common epithelial proliferation that presents in

life period





What's the diagnosis?

Skin or conj? Skin

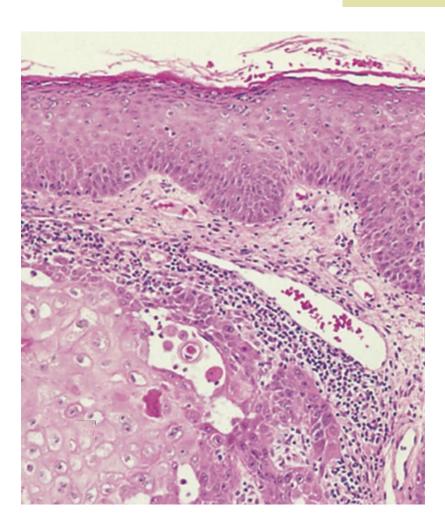
What are we supposed to notice? A few things:

- --The overall configuration is dome -shaped
- --The epi is hyperplastic, and looks benign
- --The lesion has a number of cysts containing concentrically laminated collections of surface keratin

Here is a close-up wherein the concentric lamination is easier to appreciate (both *asterisk* and *arrows*)

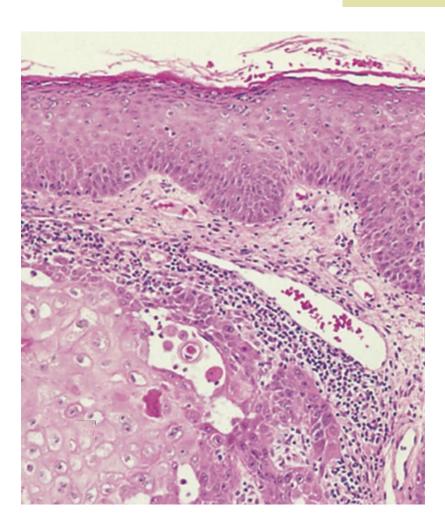
If you encounter a dome-shaped lesion containing such cysts, one diagnosis should come to mind:

Seborrheic keratosis is a common epithelial proliferation that presents in middle age .



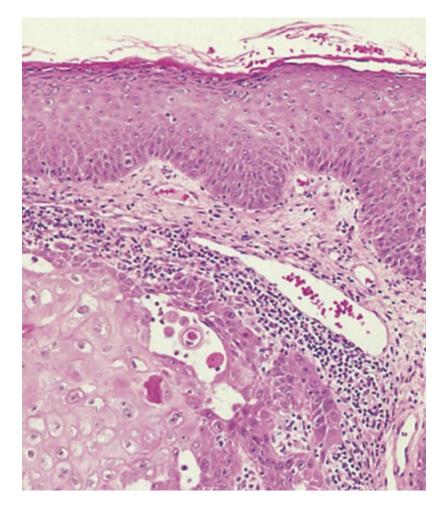
Skin, or conj?





Skin, or conj? Skin

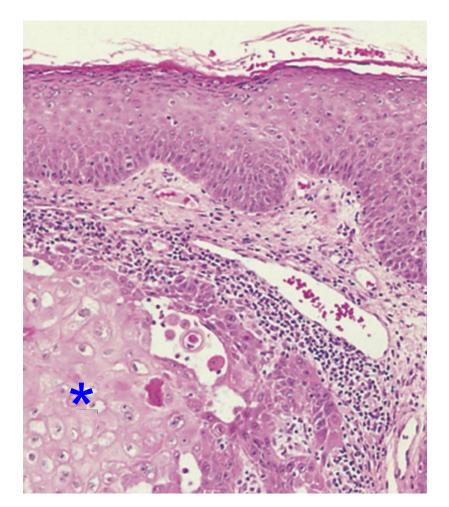






Skin, or conj? Skin

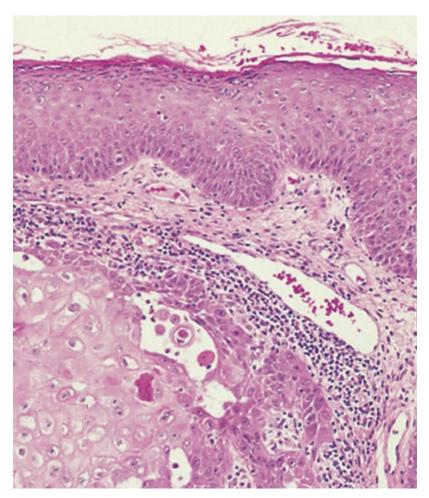
What's going on here? Note that the epithelium is quite thickened and rather chaotic in appearance. The dermis looks hinky as well.



326

Skin, or conj? Skin

What's going on here? Note that the epithelium is quite thickened and rather chaotic in appearance. The dermis looks hinky as well. But what really stands out is **this** large area of gnarly-looking squamous cells located well into the dermis.

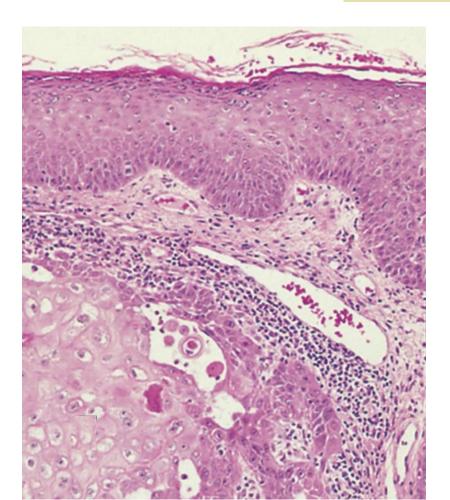


What's the diagnosis?



Skin, or conj? Skin

What's going on here? Note that the epithelium is quite thickened and rather chaotic in appearance. The dermis looks hinky as well. But what really stands out is **this** large area of gnarly-looking squamous cells located well into the dermis. Needless to say (I hope), the presence of severely atypical squamous cells in the dermis—ie, having apparently broken through their basement membrane—is strongly suggestive of one diagnosis:





What's going on here? Note that the epithelium is quite thickened and rather chaotic in appearance. The dermis looks hinky as well. But what really stands out is this large area of gnarly-looking squamous cells located well into the dermis. Needless to say (I hope), the presence of severely atypical squamous cells in the dermis—ie, having apparently broken through their basement membrane—is strongly suggestive of one diagnosis:

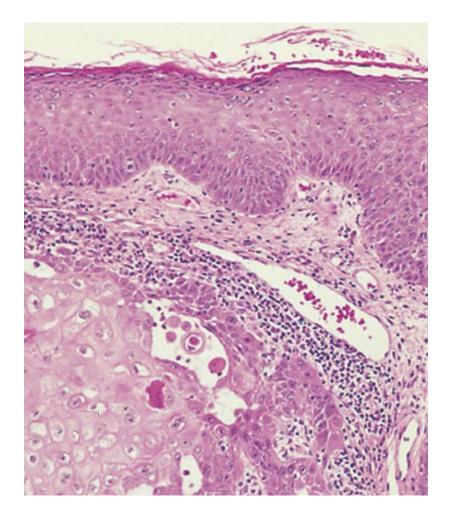
What's the diagnosis?

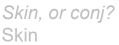
Squamous cell carcinoma (SCC) is far less v more



common than basal cell carcinoma (BCC) in the lids.





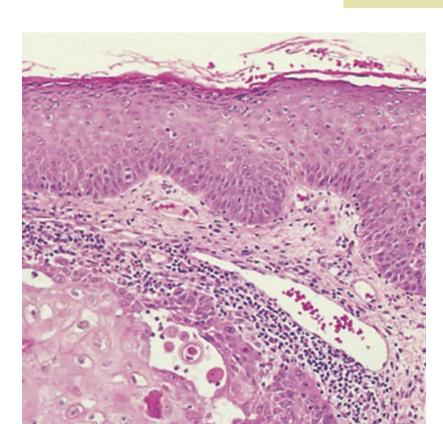


What's going on here? Note that the epithelium is quite thickened and rather chaotic in appearance. The dermis looks hinky as well. But what really stands out is **this** large area of gnarly-looking squamous cells located well into the dermis. Needless to say (I hope), the presence of severely atypical squamous cells in the dermis—ie, having apparently broken through their basement membrane—is strongly suggestive of one diagnosis:

What's the diagnosis?

Squamous cell carcinoma (SCC) is far less common than basal cell carcinoma (BCC) in the lids.





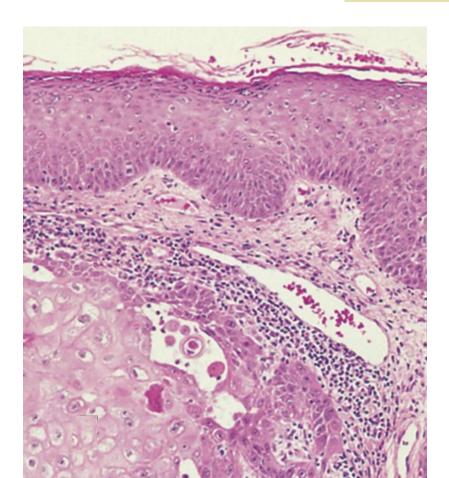


Skin, or conj? Skin

What's going on here? Note that the epithelium is quite thickened and rather chaotic in appearance. The dermis looks hinky as well. But what really stands out is **this** large area of gnarly-looking squamous cells located well into the dermis. Needless to say (I hope), the presence of severely atypical squamous cells in the dermis—ie, having apparently broken through their basement membrane—is strongly suggestive of one diagnosis:

What's the diagnosis?

Squamous cell carcinoma (SCC) is far less common than basal cell carcinoma (BCC) in the lids. Like BCC, it has a proclivity for the lower vupper lid.





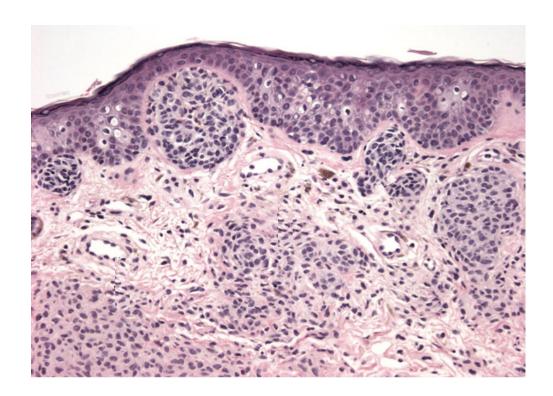
Skin, or conj? Skin

What's going on here? Note that the epithelium is quite thickened and rather chaotic in appearance. The dermis looks hinky as well. But what really stands out is **this** large area of gnarly-looking squamous cells located well into the dermis. Needless to say (I hope), the presence of severely atypical squamous cells in the dermis—ie, having apparently broken through their basement membrane—is strongly suggestive of one diagnosis:

What's the diagnosis?

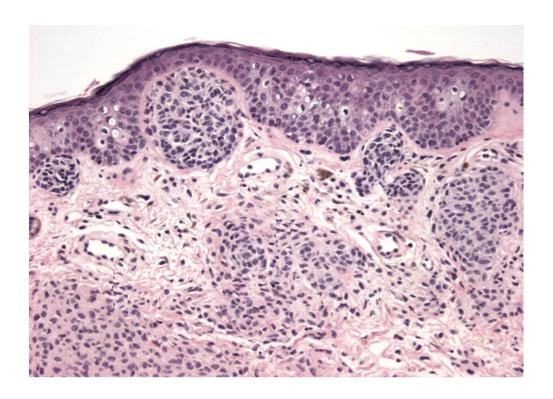
Squamous cell carcinoma (SCC) is far less common than basal cell carcinoma (BCC) in the lids. Like BCC, it has a proclivity for the lower lid.





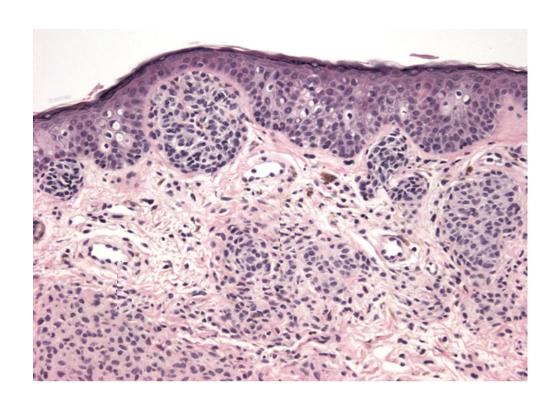
Skin/conj?





*Skin/conj?*Skin

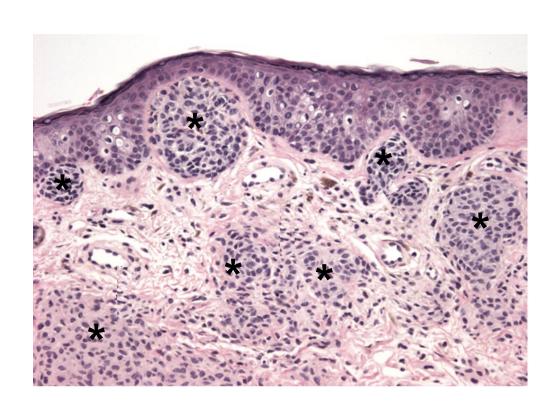




Skin/conj? Skin

There's a field mark that nails the diagnosis. What is it?



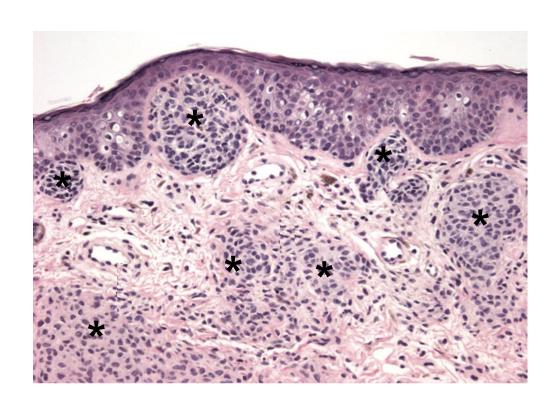


*Skin/conj?*Skin

There's a field mark that nails the diagnosis. What is it?

These tight clusters of cell type are called (aka).



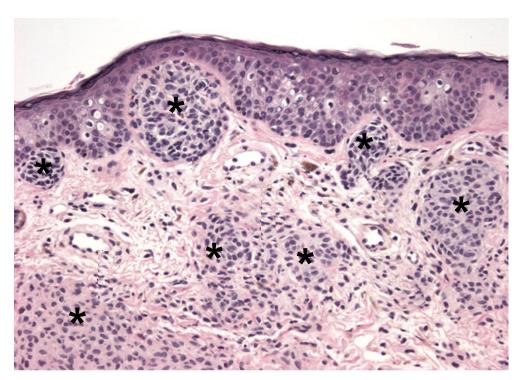


Skin/conj? Skin

There's a field mark that nails the diagnosis. What is it?

These tight clusters of melanocytes are called nests (aka *theques*).





What's the diagnosis?

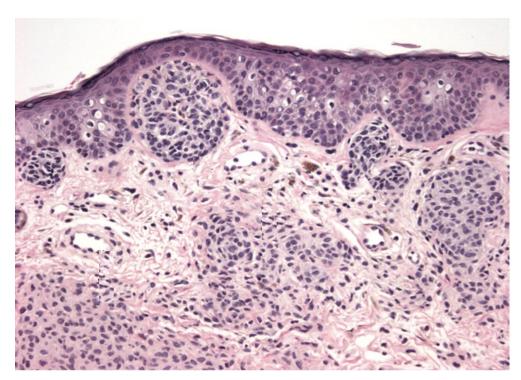
Skin/conj? Skin

There's a field mark that nails the diagnosis. What is it?

These tight clusters of melanocytes are called nests (aka *theques*).

When you encounter lid skin with melanocytes in nests, one condition should come to mind:





Skin/conj? Skin

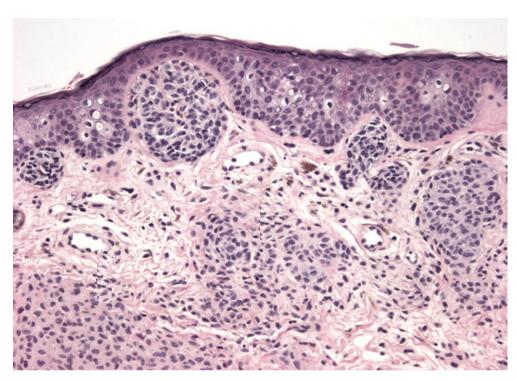
There's a field mark that nails the diagnosis. What is it?

These tight clusters of melanocytes are called nests (aka *theques*). When you encounter lid skin with melanocytes in nests, one condition should come to mind:

What's the diagnosis?

Melanocytic nevi of the lid can be congenital, or arise later in life.





Skin/conj? Skin

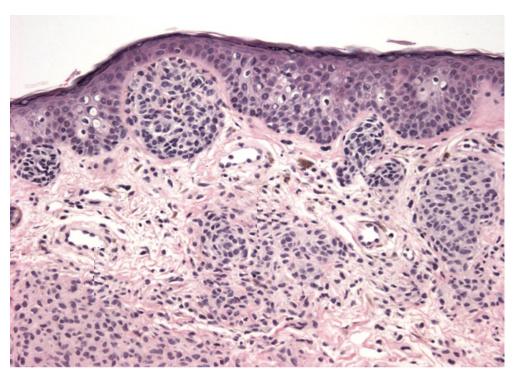
There's a field mark that nails the diagnosis. What is it?

These tight clusters of melanocytes are called nests (aka *theques*). When you encounter lid skin with melanocytes in nests, one condition should come to mind:

What's the diagnosis?

Melanocytic nevi of the lid can be congenital, or arise later in life. As with conj nevi, they evolve through a series of histologic conformations, from (in order) to to





Skin/conj? Skin

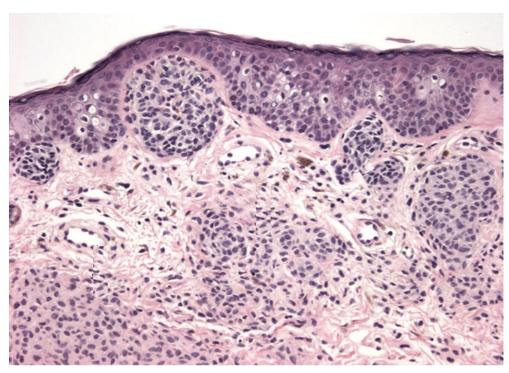
There's a field mark that nails the diagnosis. What is it?

These tight clusters of melanocytes are called nests (aka *theques*). When you encounter lid skin with melanocytes in nests, one condition should come to mind:

What's the diagnosis?

Melanocytic nevi of the lid can be congenital, or arise later in life. As with conj nevi, they evolve through a series of histologic conformations, from (in order) junctional to compound to intradermal.





Skin/conj? Skin

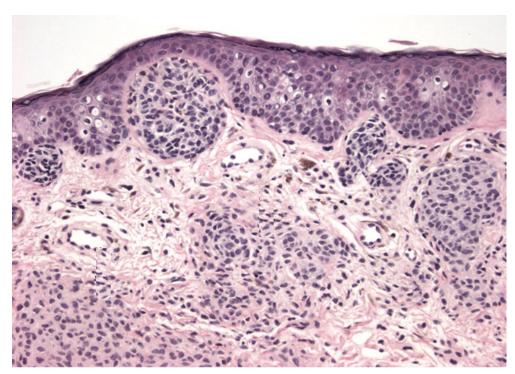
There's a field mark that nails the diagnosis. What is it?

These tight clusters of melanocytes are called nests (aka *theques*). When you encounter lid skin with melanocytes in nests, one condition should come to mind:

What's the diagnosis?

Melanocytic nevi of the lid can be congenital, or arise later in life. As with conj nevi, they evolve through a series of histologic conformations, from (in order) junctional to compound to intradermal. Most cutaneous nevi of the lid have malignant potential





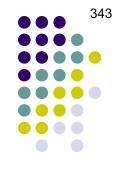
Skin/conj? Skin

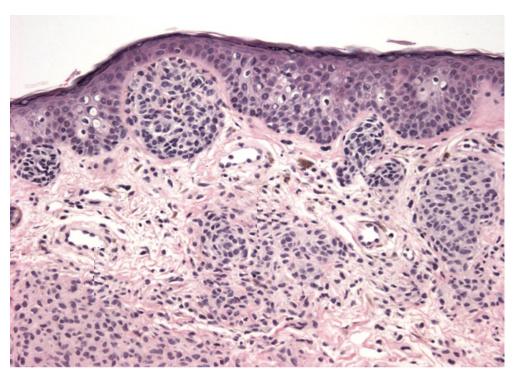
There's a field mark that nails the diagnosis. What is it?

These tight clusters of melanocytes are called nests (aka *theques*). When you encounter lid skin with melanocytes in nests, one condition should come to mind:

What's the diagnosis?

Melanocytic nevi of the lid can be congenital, or arise later in life. As with conj nevi, they evolve through a series of histologic conformations, from (in order) junctional to compound to intradermal. Most cutaneous nevi of the lid have low malignant potential





Skin/conj? Skin

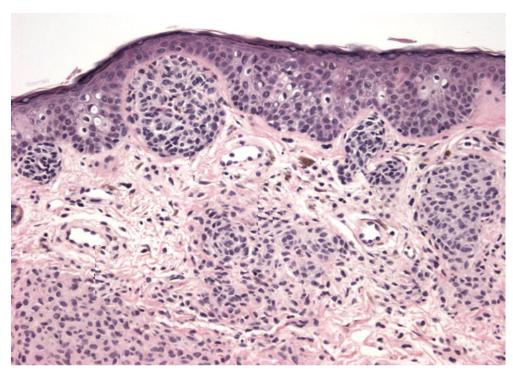
There's a field mark that nails the diagnosis. What is it?

These tight clusters of melanocytes are called nests (aka *theques*). When you encounter lid skin with melanocytes in nests, one condition should come to mind:

What's the diagnosis?

Melanocytic nevi of the lid can be congenital, or arise later in life. As with conj nevi, they evolve through a series of histologic conformations, from (in order) junctional to compound to intradermal. Most cutaneous nevi of the lid have low malignant potential; however, specific term nevi (those larger than # cm; those with irregular or) are at higher risk and should be monitored closely.





Skin/conj? Skin

There's a field mark that nails the diagnosis. What is it?

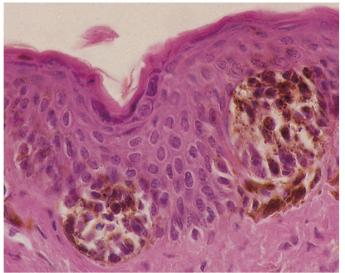
These tight clusters of melanocytes are called nests (aka *theques*). When you encounter lid skin with melanocytes in nests, one condition should come to mind:

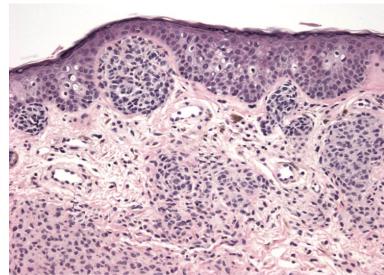
What's the diagnosis?

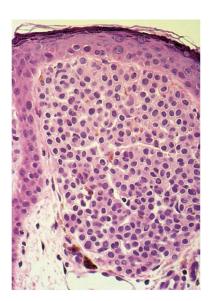
Melanocytic nevi of the lid can be congenital, or arise later in life. As with conj nevi, they evolve through a series of histologic conformations, from (in order) junctional to compound to intradermal. Most cutaneous nevi of the lid have low malignant potential; however, *dysplastic* nevi (those larger than 0.5 cm; those with irregular margins or pigmentation) are at higher risk and should be monitored closely.





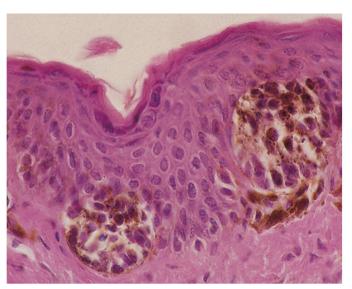


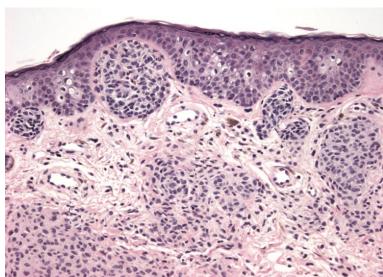


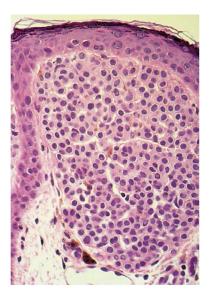


All three above are melanocytic nevi of the lid. In what important way do they differ from one another?



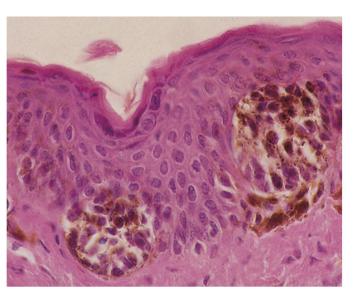


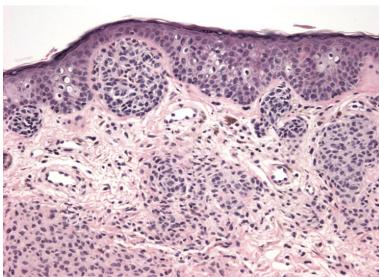


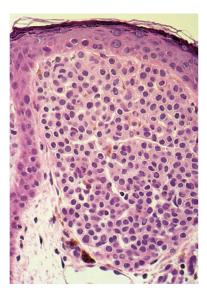


All three above are melanocytic nevi of the lid. In what important way do they differ from one another? Each represents a different stage in a nevus's conformational 'life cycle.'







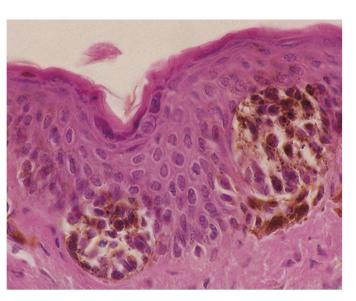


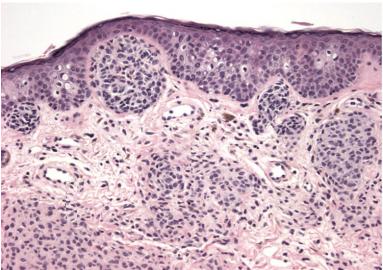
?

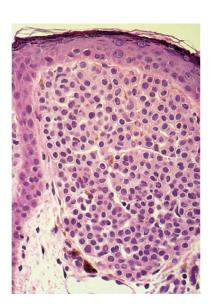
All three above are melanocytic nevi of the lid. In what important way do they differ from one another?

Each represents a different stage in a nevus's conformational 'life cycle.' In the first, the nests are confined to the dermal-epidermal junction and is therefore a nevus.





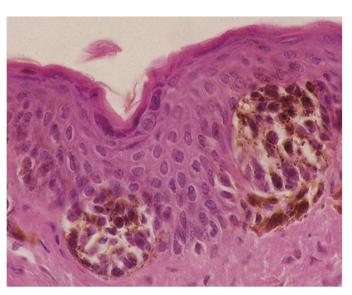


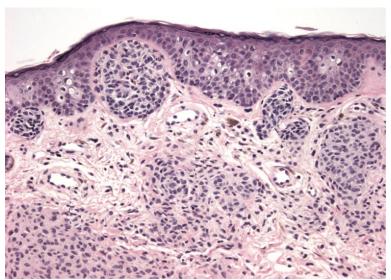


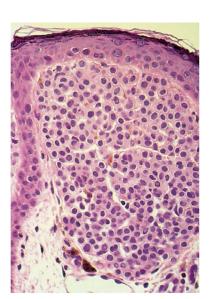
Junctional nevus

All three above are melanocytic nevi of the lid. In what important way do they differ from one another? Each represents a different stage in a nevus's conformational 'life cycle.' In the first, the nests are confined to the dermal-epidermal junction and is therefore a junctional nevus.







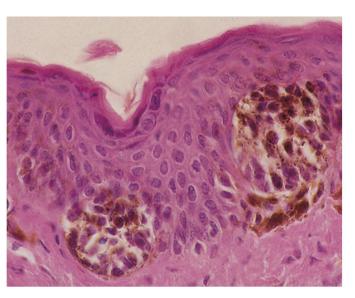


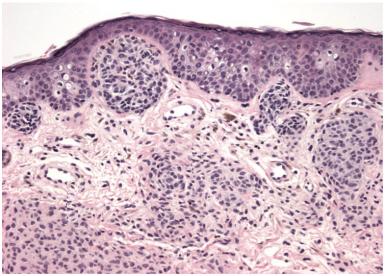
Junctional nevus

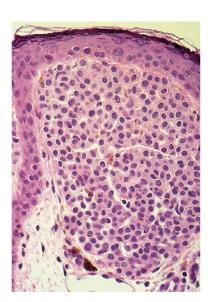
1

All three above are melanocytic nevi of the lid. In what important way do they differ from one another? Each represents a different stage in a nevus's conformational 'life cycle.' In the first, the nests are confined to the dermal-epidermal junction and is therefore a junctional nevus. In the second, the nests are found both at the dermal-epidermal junction as well as in the dermis itself; thus, it is a nevus.







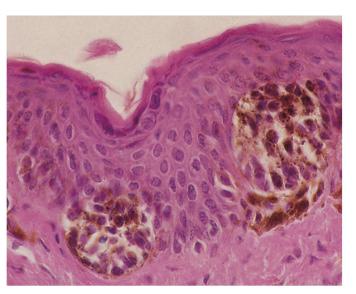


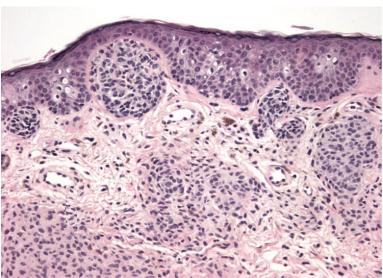
Junctional nevus

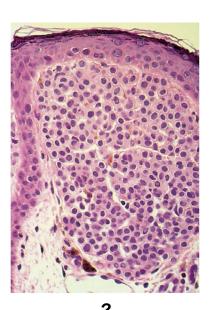
Compound nevus

All three above are melanocytic nevi of the lid. In what important way do they differ from one another? Each represents a different stage in a nevus's conformational 'life cycle.' In the first, the nests are confined to the dermal-epidermal junction and is therefore a junctional nevus. In the second, the nests are found both at the dermal-epidermal junction as well as in the dermis itself; thus, it is a compound nevus.









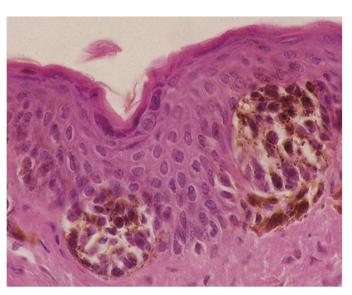
Junctional nevus

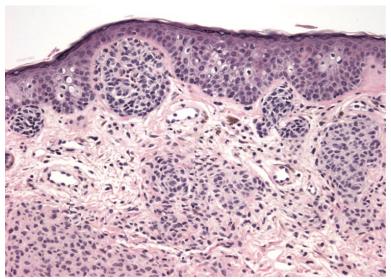
Compound nevus

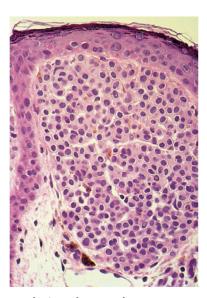
All three above are melanocytic nevi of the lid. In what important way do they differ from one another?

Each represents a different stage in a nevus's conformational 'life cycle.' In the first, the nests are confined to the dermal-epidermal junction and is therefore a junctional nevus. In the second, the nests are found both at the dermal-epidermal junction as well as in the dermis itself; thus, it is a compound nevus. In the last nevus the nest is confined to the dermis, and it therefore is an inevus.









Junctional nevus

Compound nevus

Intradermal nevus

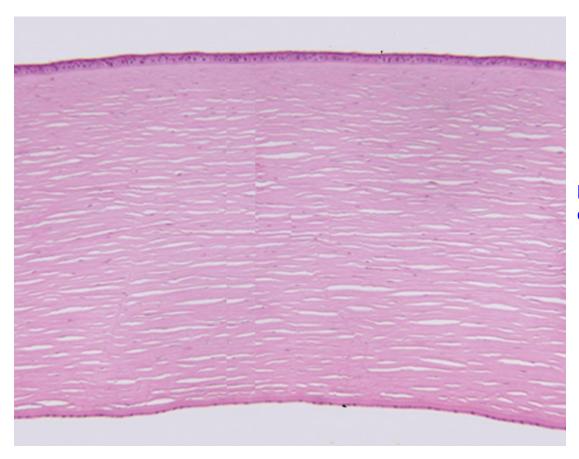
All three above are melanocytic nevi of the lid. In what important way do they differ from one another? Each represents a different stage in a nevus's conformational 'life cycle.' In the first, the nests are confined to the dermal-epidermal junction and is therefore a junctional nevus. In the second, the nests are found both at the dermal-epidermal junction as well as in the dermis itself; thus, it is a compound nevus. In the last nevus the nest is confined to the dermis, and it therefore is an intradermal nevus.



Cornea

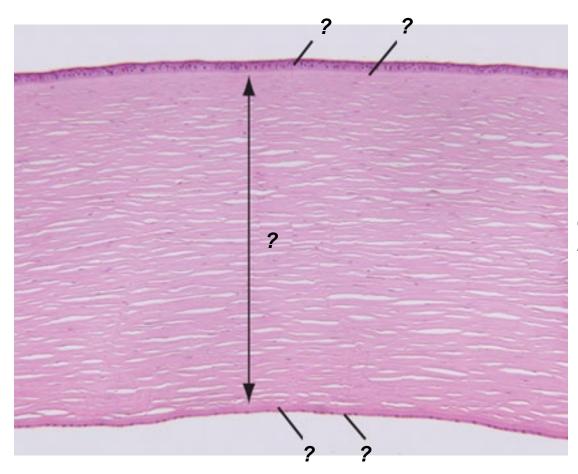
Cornea photomicrographs are a high-yield topic. Study them.





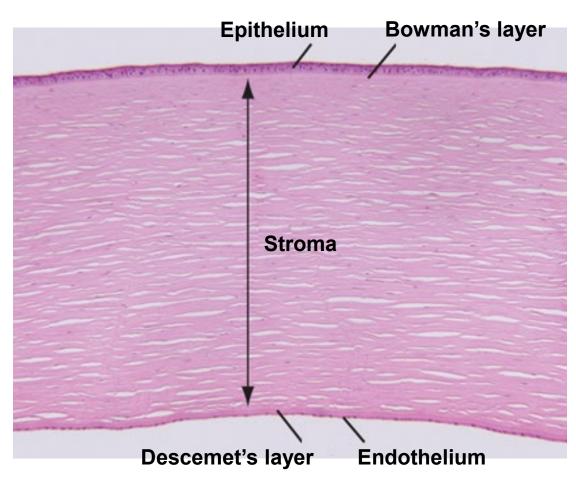
Let's spend a few minutes reviewing normal corneal histology.





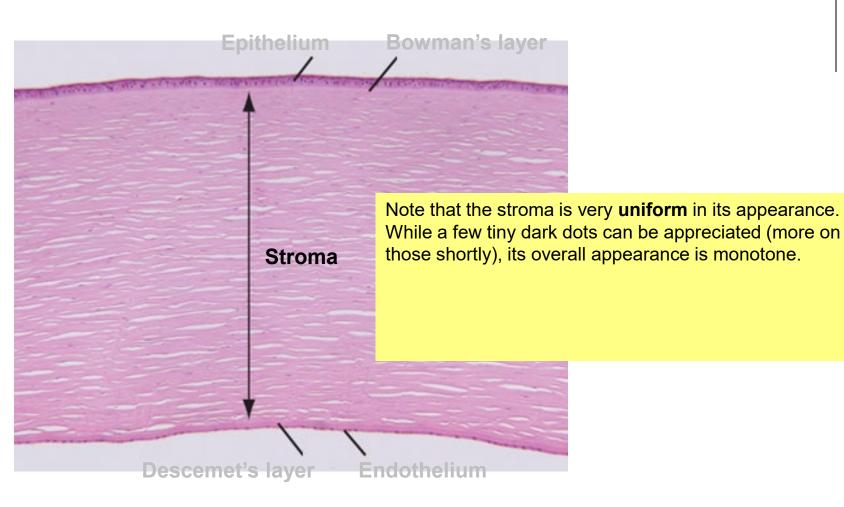
Let's spend a few minutes reviewing normal corneal histology. First, *ID the five basic layers of the cornea:*





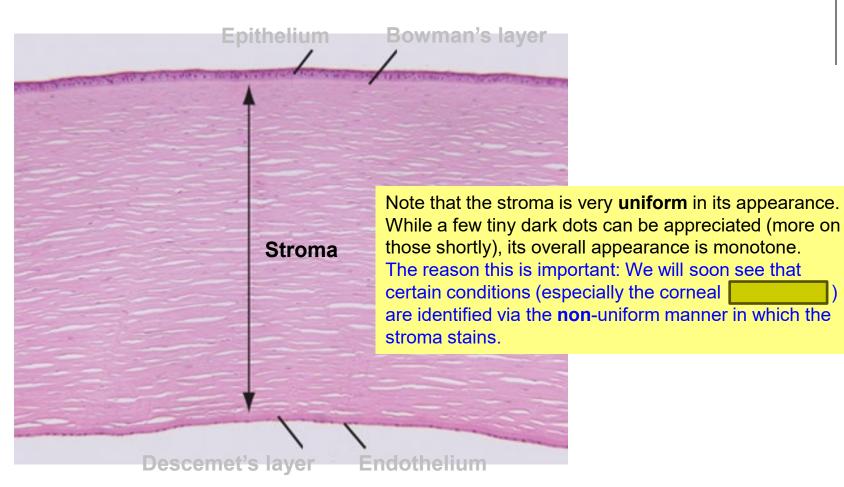
Let's spend a few minutes reviewing normal corneal histology. First, *ID the five basic layers of the cornea:*





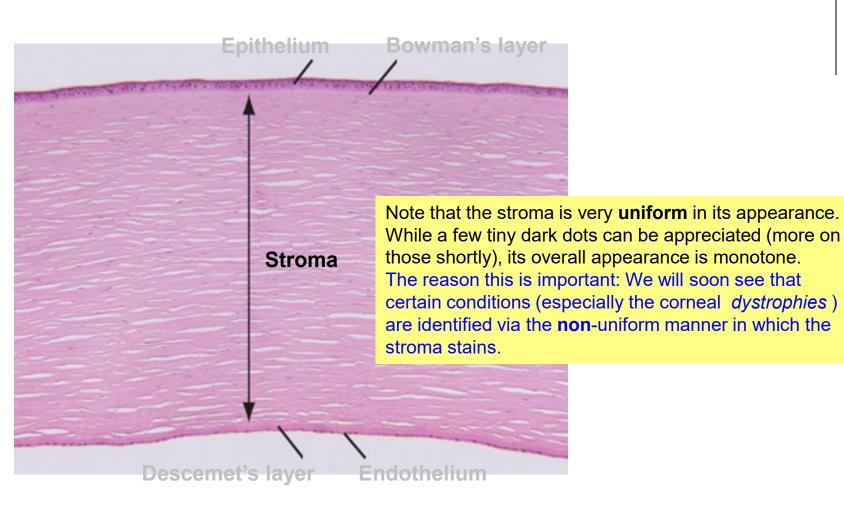
ing normal basic





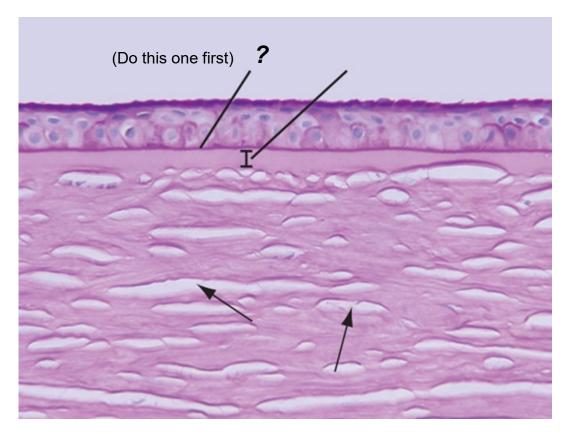
ing normal *basic*





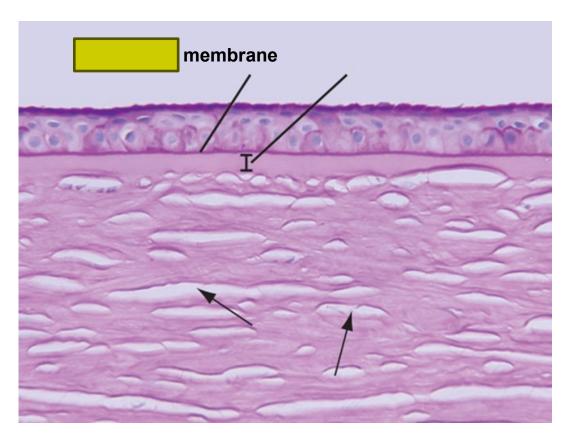
ing normal *basic*



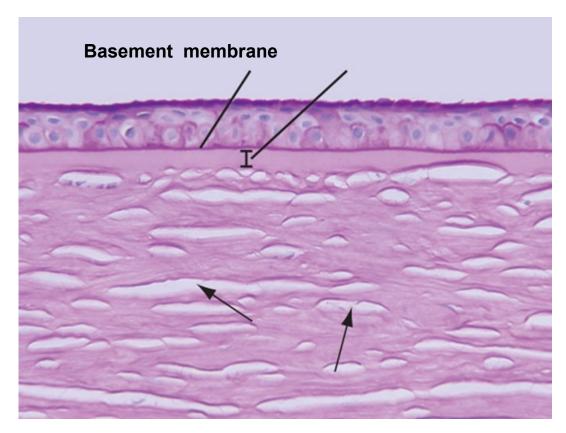


Now on with the review. This image is drilling down on the anterior cornea. *ID the indicated structures*:

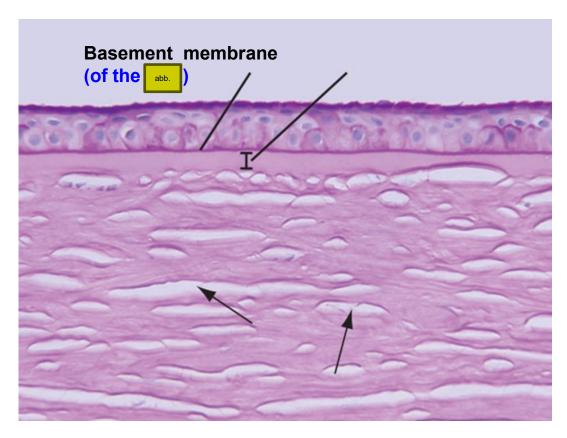




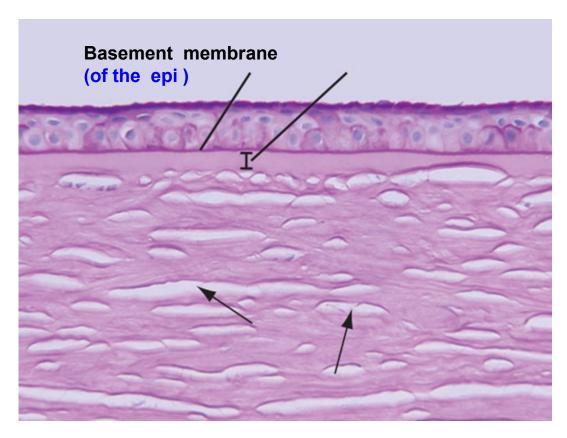




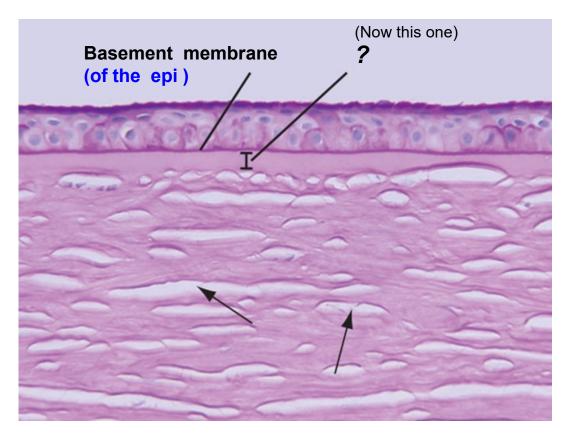




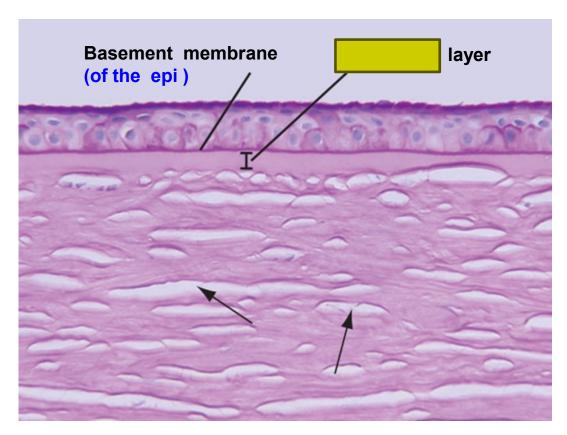




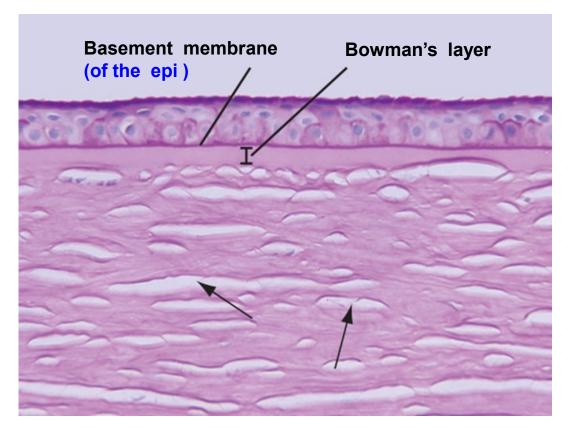




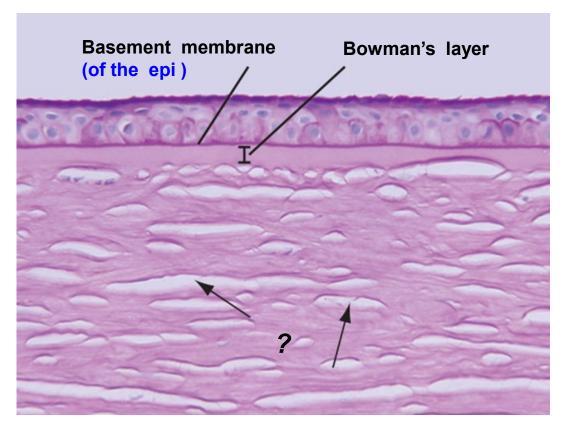








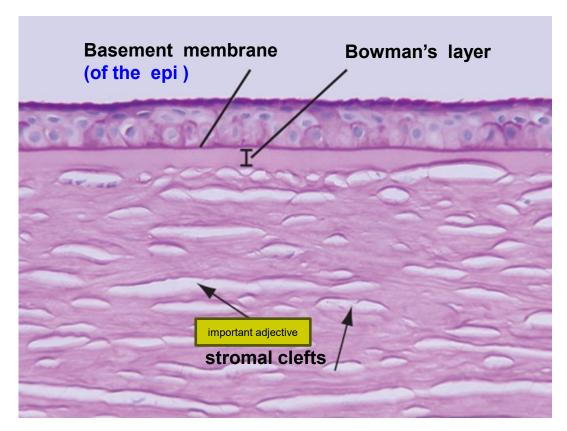




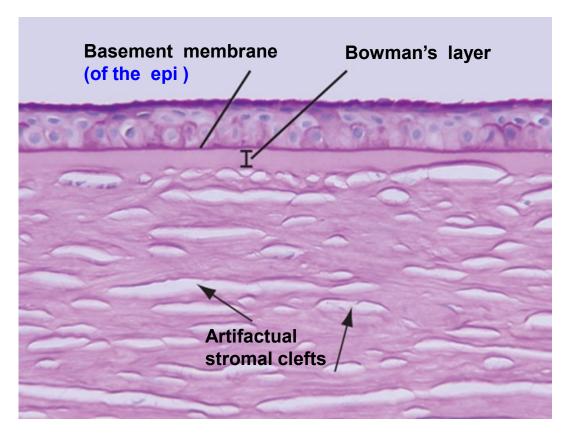
Now on with the review. This image is drilling down on the anterior cornea. *ID the indicated structures:*

(Note: Both *arrows* are pointing to examples of the same issue of interest)

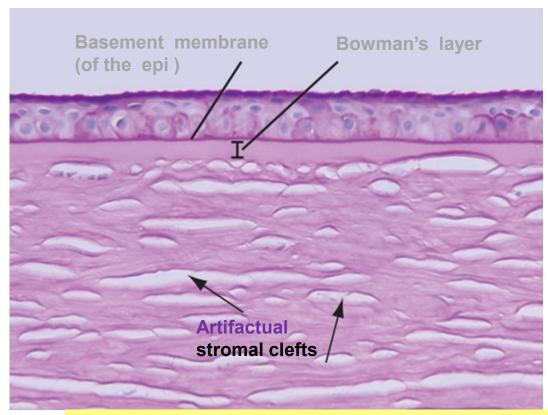








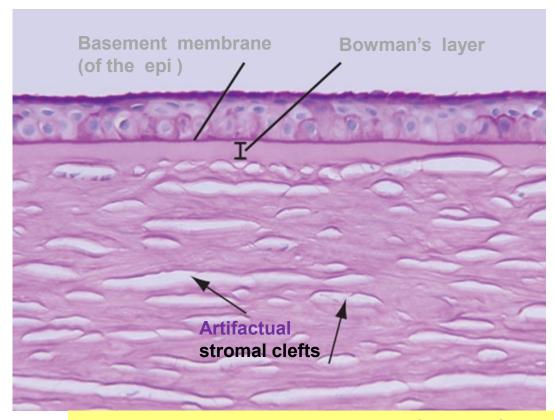




Now on with the review. This image is drilling down on the anterior cornea. *ID the indicated structures:*

What does it mean to say the stromal clefts are artifactual?



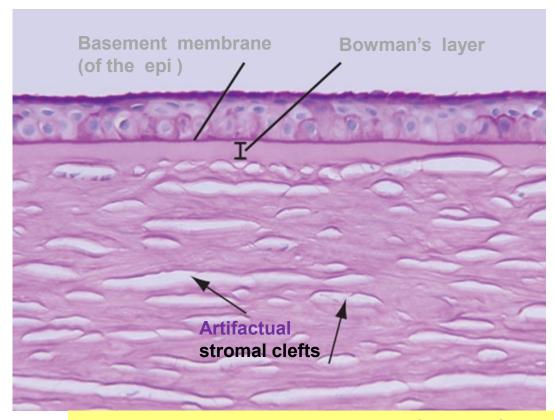


Now on with the review. This image is drilling down on the anterior cornea. *ID the indicated structures:*

What does it mean to say the stromal clefts are artifactual?

It means they arise during tissue prep, ie, are not a normal state of the tissue in vivo





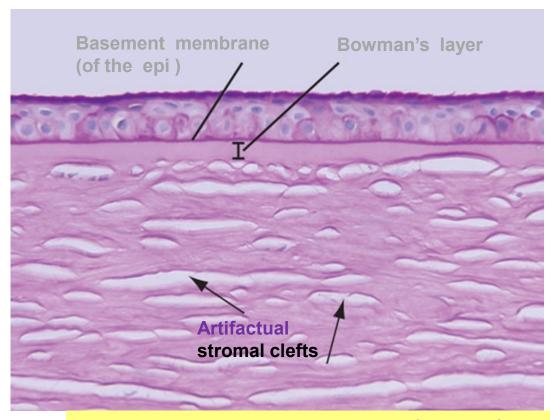
Now on with the review. This image is drilling down on the anterior cornea. *ID the indicated structures:*

What does it mean to say the stromal clefts are artifactual?

It means they arise during tissue prep, ie, are not a normal state of the tissue in vivo

OK, so the stroma contains artifactual clefts. Why should I care?





Now on with the review. This image is drilling down on the anterior cornea. *ID the indicated structures:*

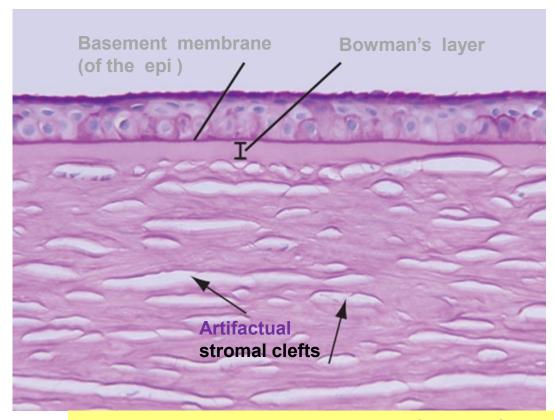
What does it mean to say the stromal clefts are artifactual?

It means they arise during tissue prep, ie, are not a normal state of the tissue in vivo

OK, so the stroma contains artifactual clefts. Why should I care?

Because this artifact is meaningful. If a portion of a cornea photomicrograph contains 'un-clefted' stroma, that portion was either or in vivo.





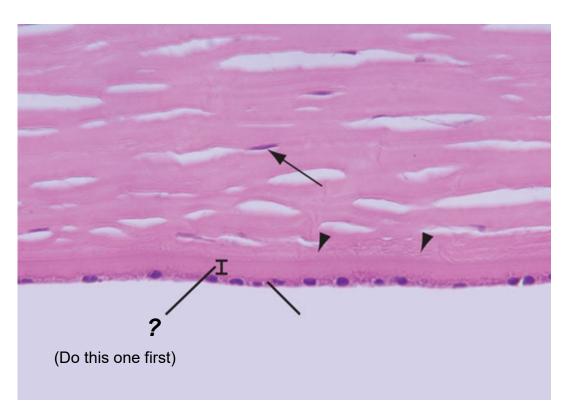
Now on with the review. This image is drilling down on the anterior cornea. *ID the indicated structures:*

What does it mean to say the stromal clefts are artifactual?
It means they arise during tissue prep, ie, are not a normal state of the tissue in vivo

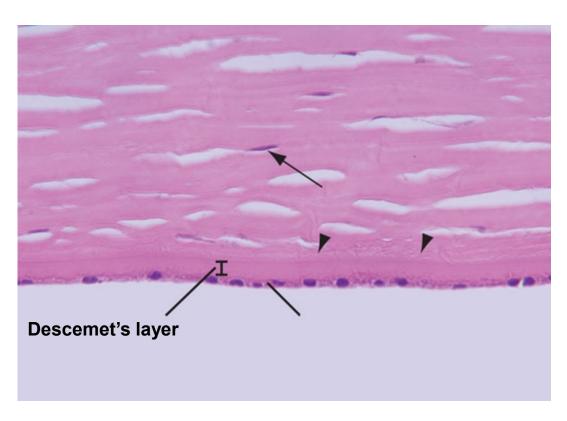
OK, so the stroma contains artifactual clefts. Why should I care?

Because this artifact is meaningful. If a portion of a cornea photomicrograph contains 'un-clefted' stroma, that portion was either scarred or edematous in vivo.

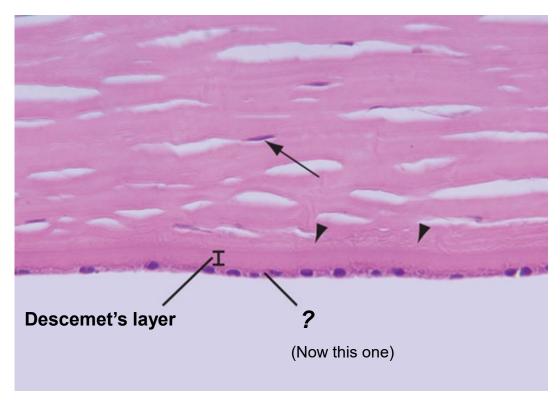




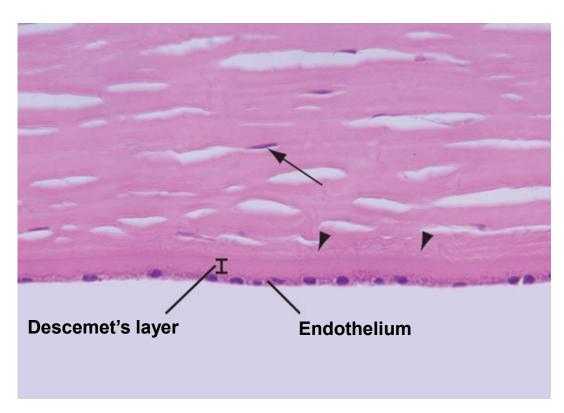




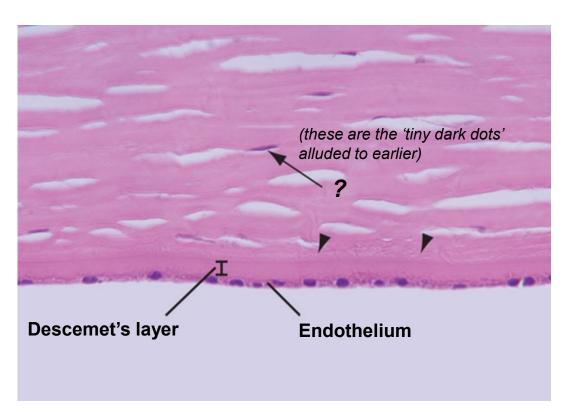




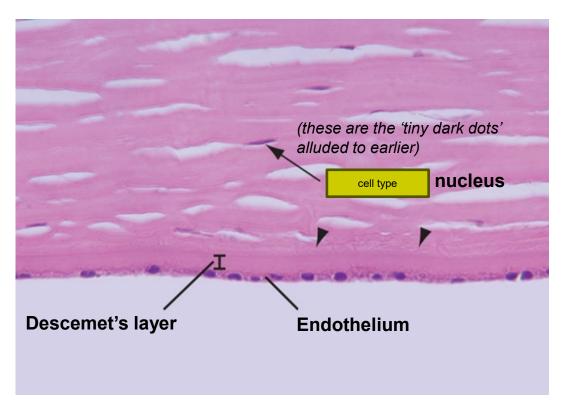




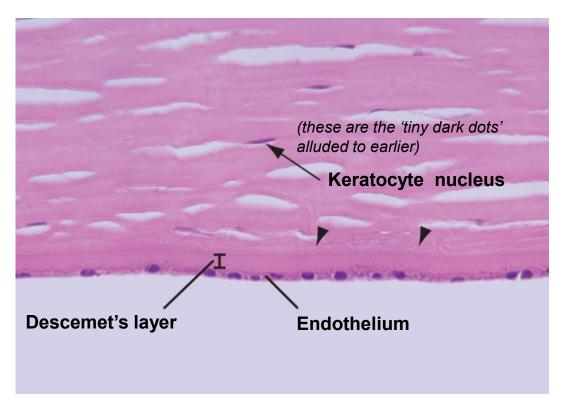




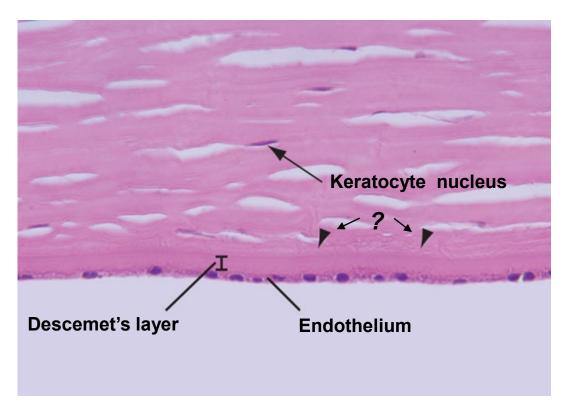




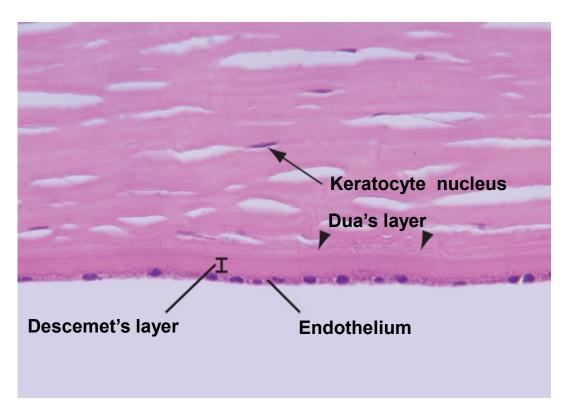




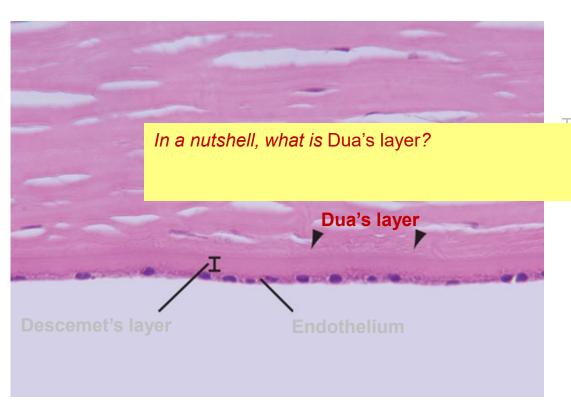




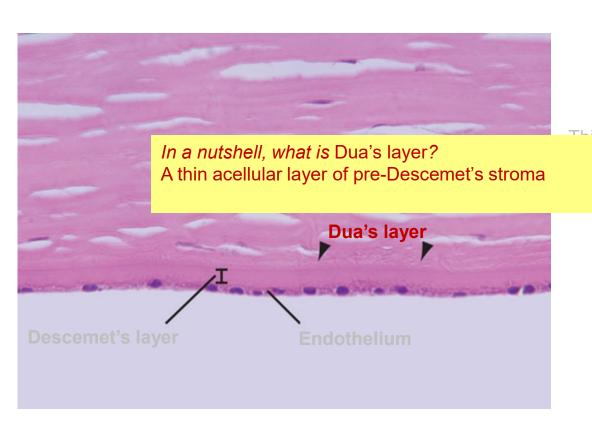




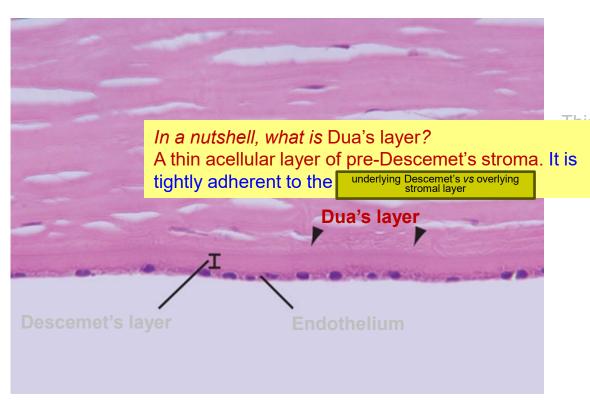




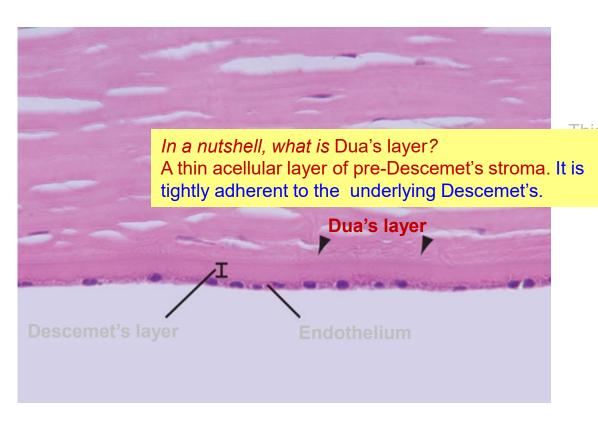










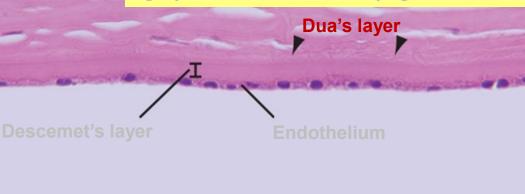


Note: Both the *Path* and *Cornea* books use *Dua layer* to refer to this portion of the cornea. That said, the term is somewhat controversial in the ophthalmic community writ large.



In a nutshell, what Dua's layer?

A thin acellular layer or pre-Descemet's stroma. It is tightly adherent to the underlying Descemet's.

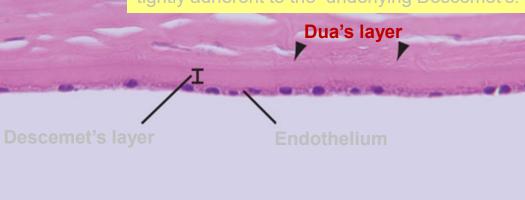


Note: Both the *Path* and *Cornea* books use *Dua layer* to refer to this portion of the cornea. That said, the term is somewhat controversial in the ophthalmic community writ large. The point being, don't be surprised if you get pushback if/when you use it, and don't necessarily look for it on an exam (ie, that portion of the cornea may be *described* rather than *named*).

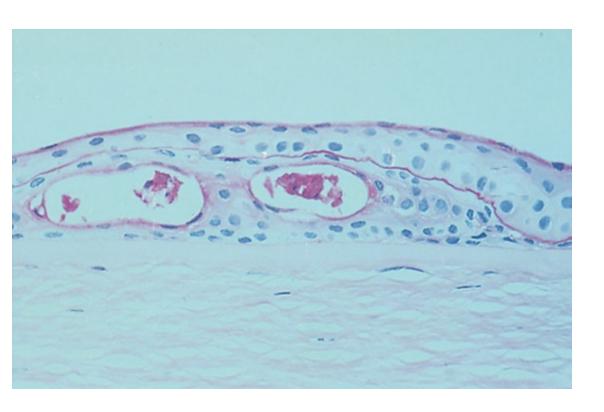


In a nutshell, what Dua's layer

A thin acellular layer of pre-Descemet's stroma. It is tightly adherent to the underlying Descemet's.

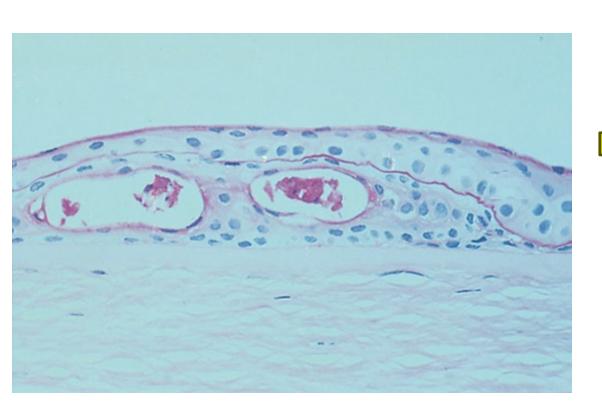






What's going on here? Before leaping to the obvious, let's take note of what looks **un**remarkable:



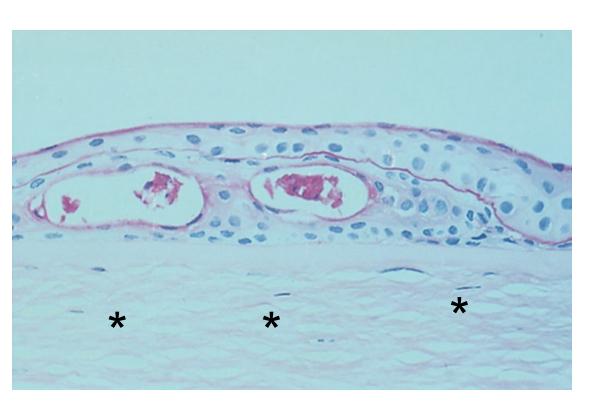


What's going on here? Before leaping to the obvious, let's take note of what looks **un**remarkable:

--The stroma contains the expected

two words

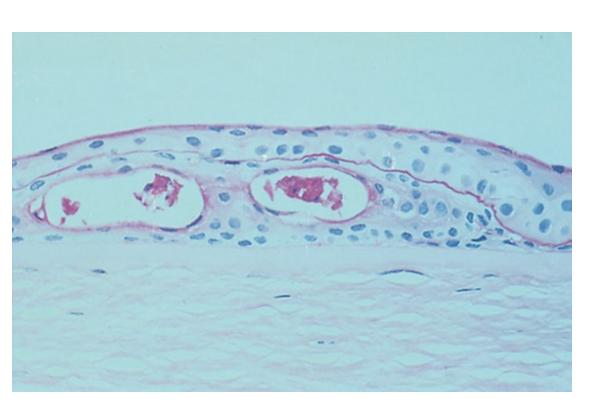




What's going on here? Before leaping to the obvious, let's take note of what looks **un**remarkable:

--The stroma contains the expected artifactual clefts



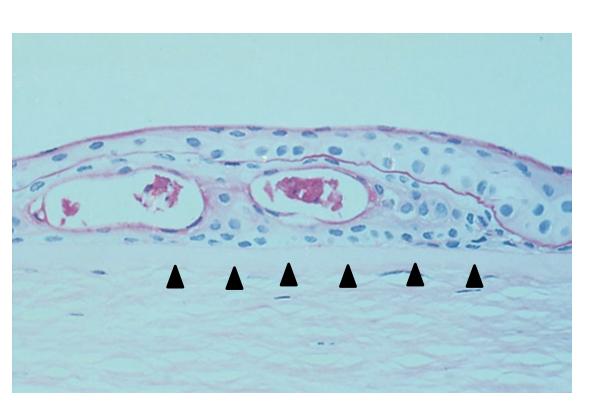


What's going on here? Before leaping to the obvious, let's take note of what looks **un**remarkable:

--The stroma contains the expected artifactual clefts; and

-- two words seems intact (albeit hard to discern with this stain).

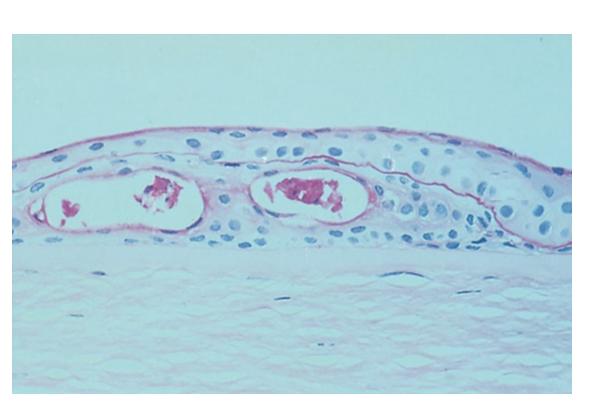




What's going on here? Before leaping to the obvious, let's take note of what looks **un**remarkable:

- --The stroma contains the expected artifactual clefts; and
- --Bowman's layer seems intact (albeit hard to discern with this stain).





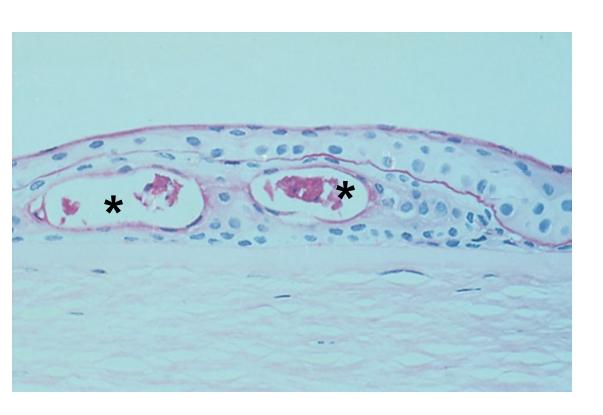
What's going on here? Before leaping to the obvious, let's take note of what looks **un**remarkable:

- --The stroma contains the expected artifactual clefts; and
- --Bowman's layer seems intact (albeit hard to discern with this stain).

Now let's talk about the obvious:

| | ا ـ ا | |
|------------|-------|--|
| The epi is | and | |
| | | |





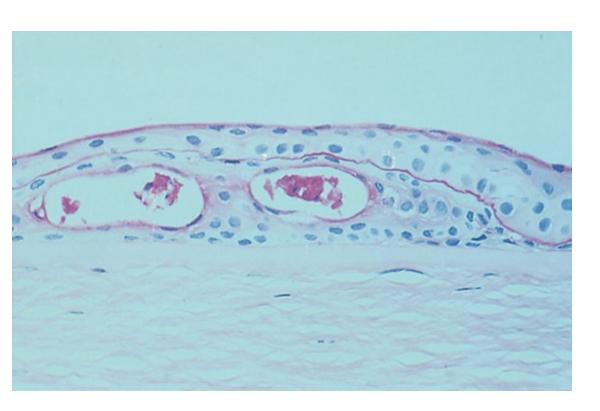
What's going on here? Before leaping to the obvious, let's take note of what looks **un**remarkable:

- --The stroma contains the expected artifactual clefts; and
- --Bowman's layer seems intact (albeit hard to discern with this stain).

Now let's talk about the obvious:

--The epi is thickened and cystic





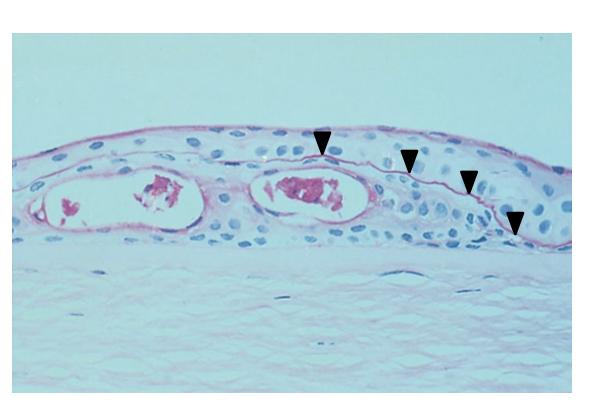
What's going on here? Before leaping to the obvious, let's take note of what looks **un**remarkable:

- --The stroma contains the expected artifactual clefts; and
- --Bowman's layer seems intact (albeit hard to discern with this stain).

Now let's talk about the obvious:

- --The epi is thickened and cystic
- --The is running up into the epithelium





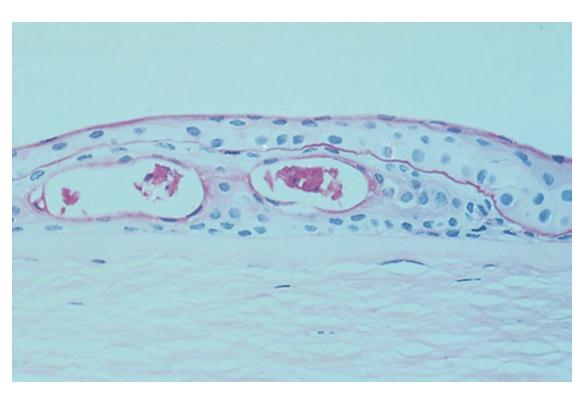
What's going on here? Before leaping to the obvious, let's take note of what looks **un**remarkable:

- --The stroma contains the expected artifactual clefts; and
- --Bowman's layer seems intact (albeit hard to discern with this stain).

Now let's talk about the obvious:

- --The epi is thickened and cystic
- --The basement membrane (BM) is running up into the epithelium.





What's the diagnosis?

What's going on here? Before leaping to the obvious, let's take note of what looks **un**remarkable:

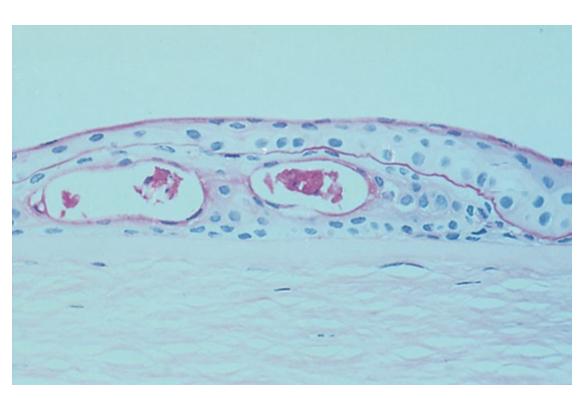
- --The stroma contains the expected artifactual clefts; and
- --Bowman's layer seems intact (albeit hard to discern with this stain).

Now let's talk about the obvious:

- --The epi is thickened and cystic
- --The basement membrane (BM) is running up into the epithelium.

A thickened epi with BM running up into it can only be one thing:





What's going on here? Before leaping to the obvious, let's take note of what looks **un**remarkable:

- --The stroma contains the expected artifactual clefts; and
- --Bowman's layer seems intact (albeit hard to discern with this stain).

Now let's talk about the obvious:
--The epi is thickened and cystic

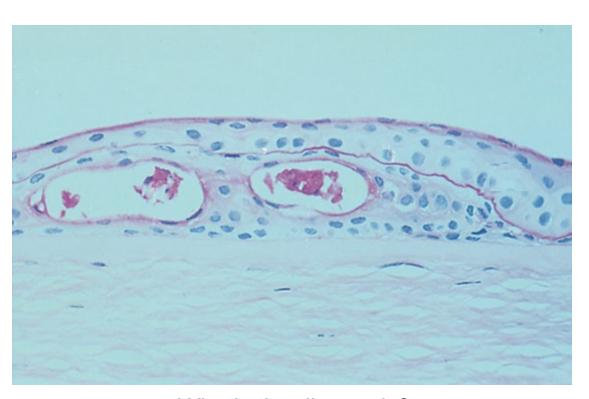
--The basement membrane (BM) is running up into the epithelium

A thickened epi with BM running up into it can only be one thing:

What's the diagnosis?

Epithelial basement membrane dystrophy (EBMD) is aka three words dystrophy on account of its appearance at the slit lamp. (It has other names as well.)





What's going on here? Before leaping to the obvious, let's take note of what looks **un**remarkable:

- --The stroma contains the expected artifactual clefts; and
- --Bowman's layer seems intact (albeit hard to discern with this stain).

Now let's talk about the obvious:

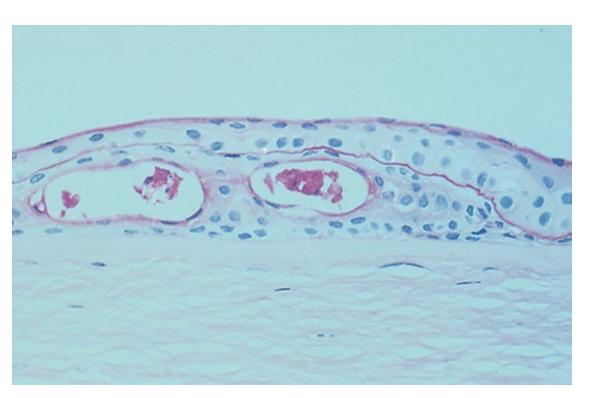
- --The epi is thickened and cystic
- --The basement membrane (BM) is running up into the epithelium

A thickened epi with BM running up into it can only be one thing:

What's the diagnosis?

Epithelial basement membrane dystrophy (EBMD) is aka *map-dot-fingerprint dystrophy* on account of its appearance at the slit lamp. (It has other names as well.)





What's going on here? Before leaping to the obvious, let's take note of what looks **un**remarkable:

- --The stroma contains the expected artifactual clefts; and
- --Bowman's layer seems intact (albeit hard to discern with this stain).

Now let's talk about the obvious:

- --The epi is thickened and cystic
- --The basement membrane (BM) is running up into the epithelium

A thickened epi with BM running up into it can only be one thing:

What's the diagnosis?

Epithelial basement membrane dystrophy (EBMD) is aka *map-dot-fingerprint dystrophy* on account of its appearance at the slit lamp. (It has other names as well.) The intraepithelial insinuation of BM accounts for the woof them , whereas the (pseudo)cysts account for the the last one





What's going on here? Before leaping to the obvious, let's take note of what looks **un**remarkable:

- --The stroma contains the expected artifactual clefts; and
- --Bowman's layer seems intact (albeit hard to discern with this stain).

Now let's talk about the obvious:

- --The epi is thickened and cystic
- --The basement membrane (BM) is running up into the epithelium

A thickened epi with BM running up into it can only be one thing:

What's the diagnosis?

Epithelial basement membrane dystrophy (EBMD) is aka *map-dot-fingerprint dystrophy* on account of its appearance at the slit lamp. (It has other names as well.) The intraepithelial insinuation of BM accounts for the *maps* and *fingerprints*, whereas the (pseudo)cysts account for the *dots*.





What's going on here? Before leaping to the obvious, let's take note of what looks **un**remarkable:

- --The stroma contains the expected artifactual clefts; and
- --Bowman's layer seems intact (albeit hard to discern with this stain).

Now let's talk about the obvious:

- --The epi is thickened and cystic
- --The basement membrane (BM) is running up into the epithelium

A thickened epi with BM running up into it can only be one thing:

What's the diagnosis?

Epithelial basement membrane dystrophy (EBMD) is aka *map-dot-fingerprint dystrophy* on account of its appearance at the slit lamp. (It has other names as well.) The intraepithelial insinuation of BM accounts for the *maps* and *fingerprints*, whereas the (pseudo)cysts account for the *dots*.

Complaints are related to three words





What's going on here? Before leaping to the obvious, let's take note of what looks **un**remarkable:

- --The stroma contains the expected artifactual clefts; and
- --Bowman's layer seems intact (albeit hard to discern with this stain).

Now let's talk about the obvious:
--The epi is thickened and cystic

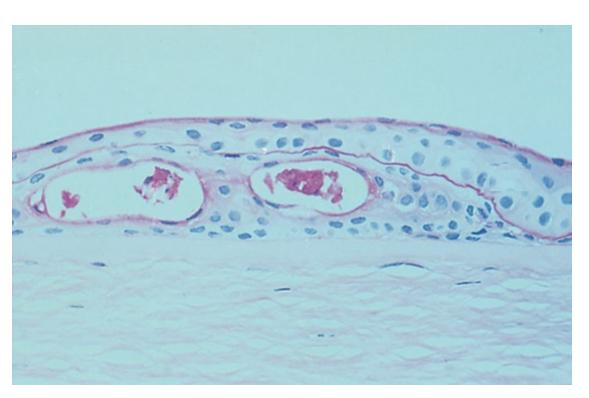
--The basement membrane (BM) is running up into the epithelium

A thickened epi with BM running up into it can only be one thing:

What's the diagnosis?

Epithelial basement membrane dystrophy (EBMD) is aka *map-dot-fingerprint dystrophy* on account of its appearance at the slit lamp. (It has other names as well.) The intraepithelial insinuation of BM accounts for the *maps* and *fingerprints*, whereas the (pseudo)cysts account for the *dots*. Complaints are related to recurrent epithelial erosions.





What's going on here? Before leaping to the obvious, let's take note of what looks **un**remarkable:

- --The stroma contains the expected artifactual clefts; and
- --Bowman's layer seems intact (albeit hard to discern with this stain).

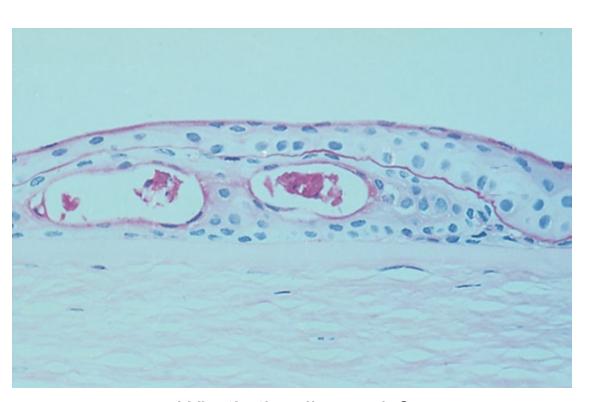
Now let's talk about the obvious:
--The epi is thickened and cystic
--The basement membrane (BM) is
running up into the epithelium

A thickened epi with BM running up into it can only be one thing:

What's the diagnosis?

Epithelial basement membrane dystrophy (EBMD) is aka *map-dot-fingerprint dystrophy* on account of its appearance at the slit lamp. (It has other names as well.) The intraepithelial insinuation of BM accounts for the *maps* and *fingerprints*, whereas the (pseudo)cysts account for the *dots*. Complaints are related to recurrent epithelial erosions. Vision typically affected.





What's going on here? Before leaping to the obvious, let's take note of what looks **un**remarkable:

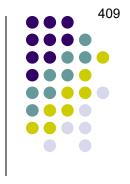
- --The stroma contains the expected artifactual clefts; and
- --Bowman's layer seems intact (albeit hard to discern with this stain).

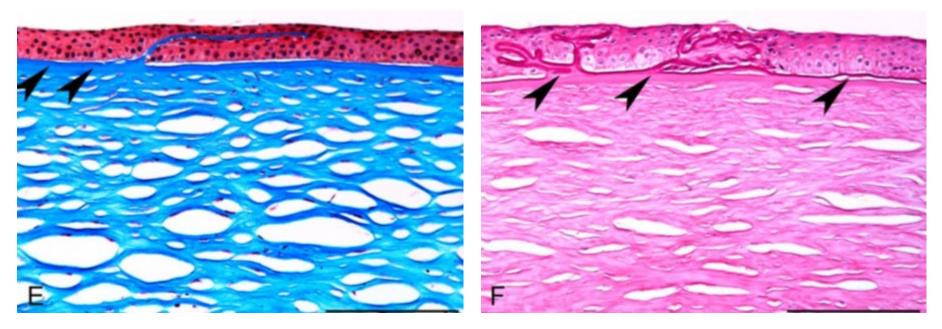
Now let's talk about the obvious:
--The epi is thickened and cystic
--The basement membrane (BM) is
running up into the epithelium

A thickened epi with BM running up into it can only be one thing:

What's the diagnosis?

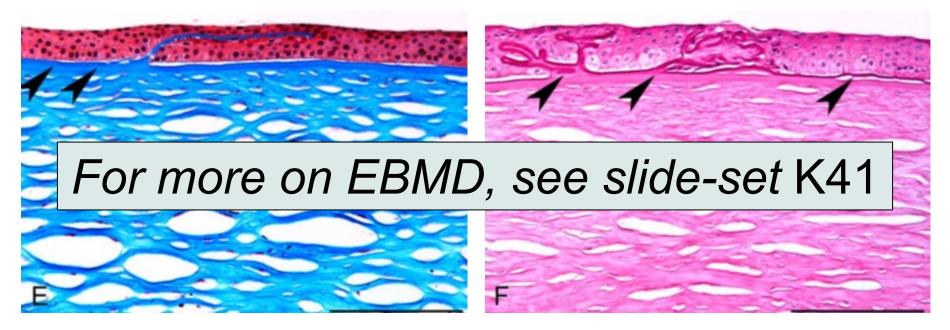
Epithelial basement membrane dystrophy (EBMD) is aka *map-dot-fingerprint dystrophy* on account of its appearance at the slit lamp. (It has other names as well.) The intraepithelial insinuation of BM accounts for the *maps* and *fingerprints*, whereas the (pseudo)cysts account for the *dots*. Complaints are related to recurrent epithelial erosions. Vision typically is affected.





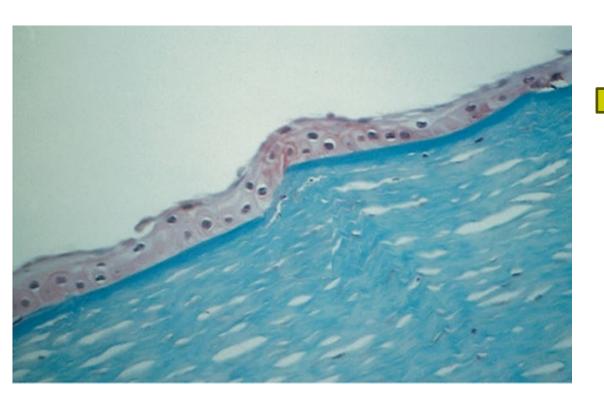
More EBMD examples wherein the BM can be better seen snaking into the epithelium





More EBMD examples wherein the BM can be better seen snaking into the epithelium

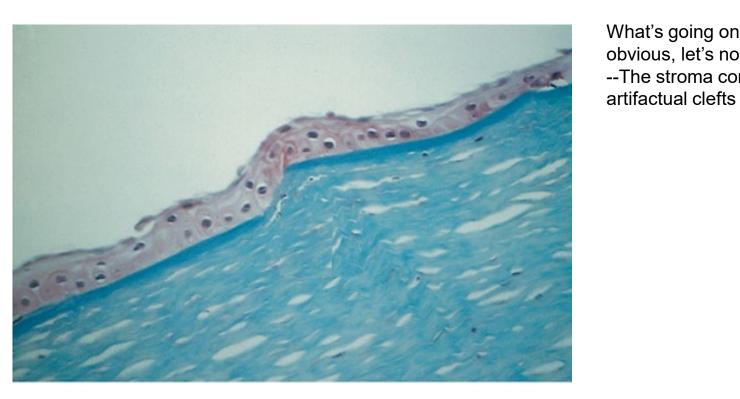




What's going on here? Again saving the obvious, let's note what looks OK:
--The stroma contains the expected

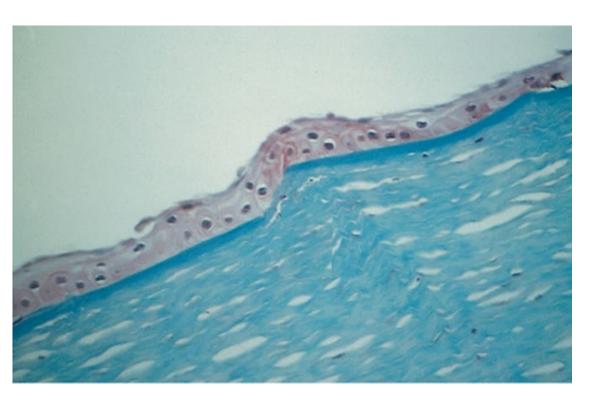
two words





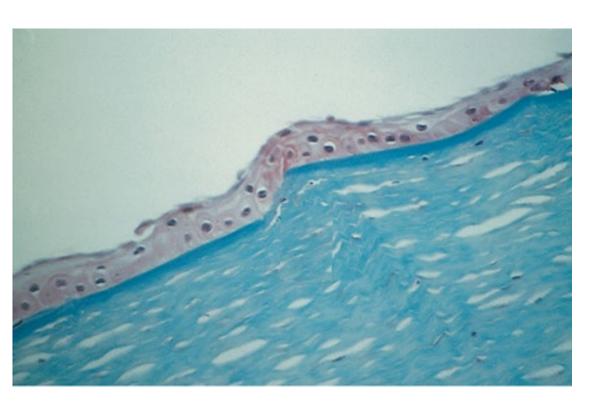
What's going on here? Again saving the obvious, let's note what looks OK:
--The stroma contains the expected





What's going on here? Again saving the obvious, let's note what looks OK:
--The stroma contains the expected artifactual clefts, and isn't picking up stain in a way that suggests it contains some sort of abnormal material.

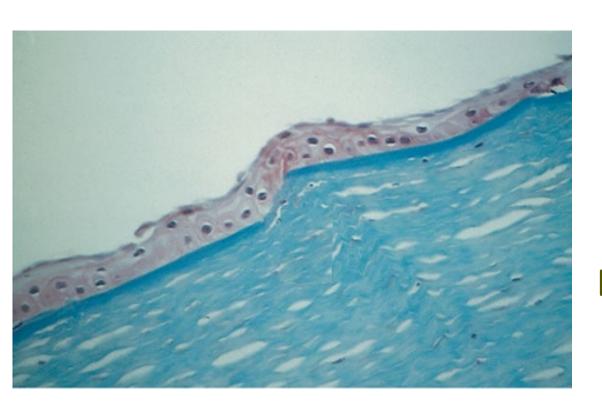




What's going on here? Again saving the obvious, let's note what looks OK:

- --The stroma contains the expected artifactual clefts, and isn't picking up stain in a way that suggests it contains some sort of abnormal material.
- --While the epithelium looks wonky, it doesn't contain discernible BM running through it.





What's going on here? Again saving the obvious, let's note what looks OK:

- --The stroma contains the expected artifactual clefts, and isn't picking up stain in a way that suggests it contains some sort of abnormal material.
- --While the epithelium looks wonky, it doesn't contain discernible BM running through it.

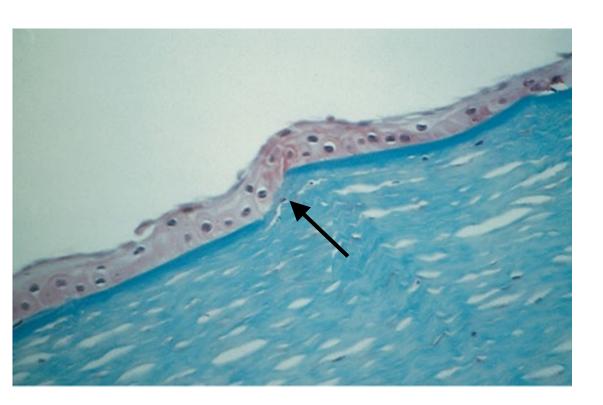
Now let's talk about the obvious:

two words

is completely disrupted

right **here**.



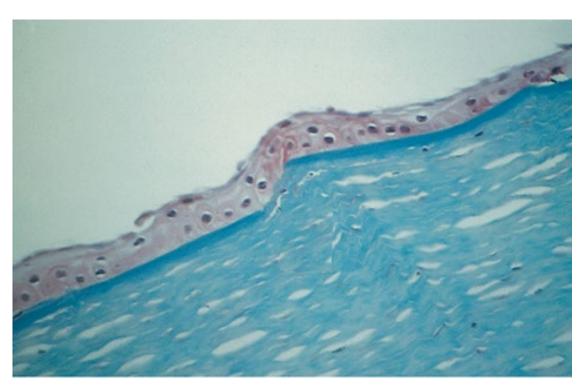


What's going on here? Again saving the obvious, let's note what looks OK:

- --The stroma contains the expected artifactual clefts, and isn't picking up stain in a way that suggests it contains some sort of abnormal material.
- --While the epithelium looks wonky, it doesn't contain discernible BM running through it.

Now let's talk about the obvious: Bowman's layer is completely disrupted right **here**.





What's the diagnosis?

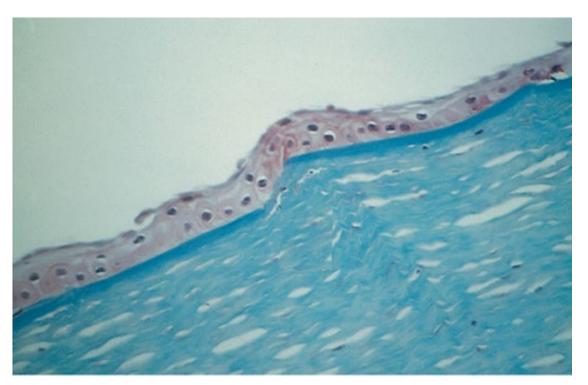
What's going on here? Again saving the obvious, let's note what looks OK:

- --The stroma contains the expected artifactual clefts, and isn't picking up stain in a way that suggests it contains some sort of abnormal material.
- --While the epithelium looks wonky, it doesn't contain discernible BM running through it.

Now let's talk about the obvious: Bowman's layer is completely disrupted right **here**.

Focal disruption of Bowman's layer in the absence of abnormal stromal staining should bring to mind one dx:





What's the diagnosis?

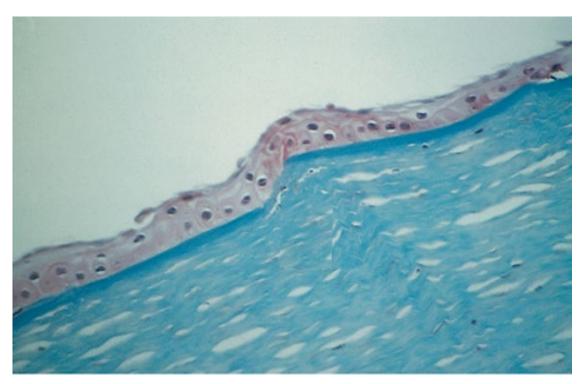
Keratoconus is an general disorder

What's going on here? Again saving the obvious, let's note what looks OK:
--The stroma contains the expected artifactual clefts, and isn't picking up stain in a way that suggests it contains some sort of abnormal material.
--While the epithelium looks wonky, it doesn't contain discernible BM running through it.

Now let's talk about the obvious: Bowman's layer is completely disrupted right **here**.

Focal disruption of Bowman's layer in the absence of abnormal stromal staining should bring to mind one dx:





What's the diagnosis?

Keratoconus is an ectatic disorder

What's going on here? Again saving the obvious, let's note what looks OK:

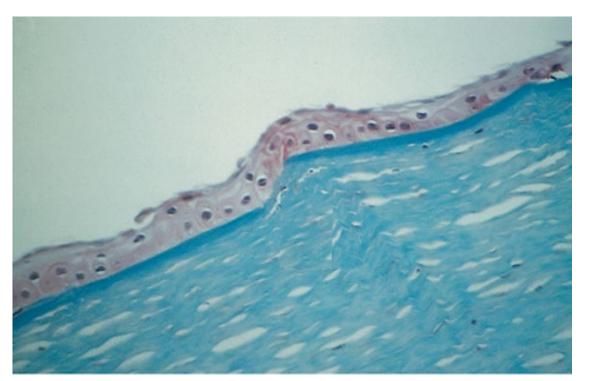
--The stroma contains the expected artifactual clefts, and isn't picking up stain in a way that suggests it contains some sort of abnormal material.

--While the epithelium looks wonky, it doesn't contain discernible BM running through it.

Now let's talk about the obvious: Bowman's layer is completely disrupted right **here**.

Focal disruption of Bowman's layer in the absence of abnormal stromal staining should bring to mind one dx:





What's the diagnosis?

What's going on here? Again saving the obvious, let's note what looks OK: --The stroma contains the expected artifactual clefts, and isn't picking up stain in a way that suggests it contains some sort of abnormal material. --While the epithelium looks wonky, it doesn't contain discernible BM running

Now let's talk about the obvious: Bowman's layer is completely disrupted right here.

Focal disruption of Bowman's layer in the absence of abnormal stromal staining should bring to mind one dx:

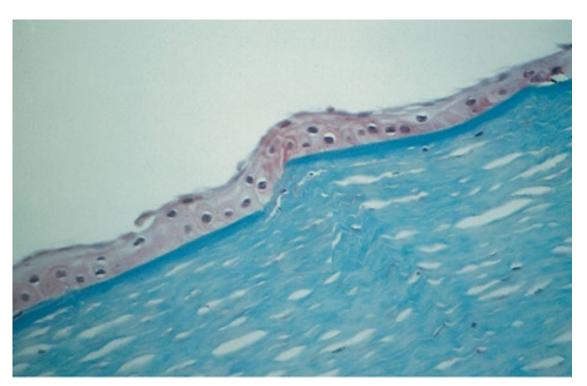
Keratoconus is an ectatic disorder characterized by progressive corneal change 1

through it.

and

change 2





What's the diagnosis?

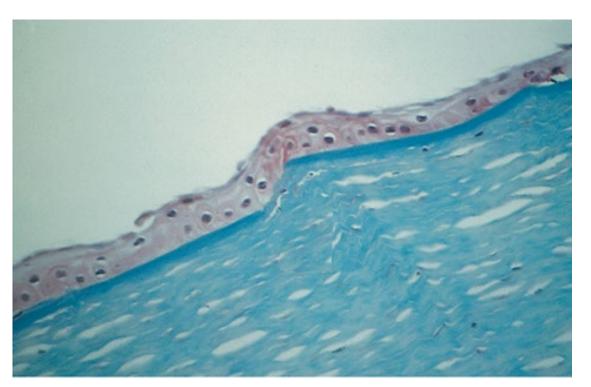
What's going on here? Again saving the obvious, let's note what looks OK:
--The stroma contains the expected artifactual clefts, and isn't picking up stain in a way that suggests it contains some sort of abnormal material.
--While the epithelium looks wonky, it doesn't contain discernible BM running through it.

Now let's talk about the obvious: Bowman's layer is completely disrupted right **here**.

Focal disruption of Bowman's layer in the absence of abnormal stromal staining should bring to mind one dx:

Keratoconus is an ectatic disorder characterized by progressive corneal thinning and protrusion





What's the diagnosis?

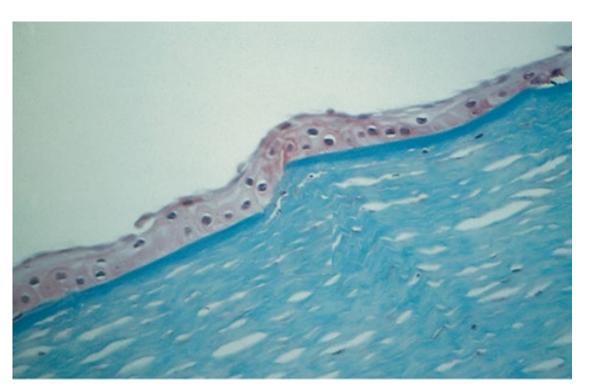
What's going on here? Again saving the obvious, let's note what looks OK:
--The stroma contains the expected artifactual clefts, and isn't picking up stain in a way that suggests it contains some sort of abnormal material.
--While the epithelium looks wonky, it doesn't contain discernible BM running through it.

Now let's talk about the obvious: Bowman's layer is completely disrupted right **here**.

Focal disruption of Bowman's layer in the absence of abnormal stromal staining should bring to mind one dx:

Keratoconus is an ectatic disorder characterized by progressive corneal thinning and protrusion of its area 1 and/or area 2 portions.





What's the diagnosis?

What's going on here? Again saving the obvious, let's note what looks OK:

--The stroma contains the expected artifactual clefts, and isn't picking up stain in a way that suggests it contains some sort of abnormal material.

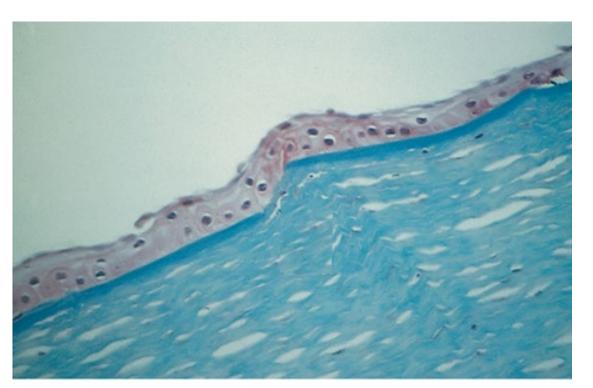
--While the epithelium looks wonky, it doesn't contain discernible BM running through it.

Now let's talk about the obvious: Bowman's layer is completely disrupted right **here**.

Focal disruption of Bowman's layer in the absence of abnormal stromal staining should bring to mind one dx:

Keratoconus is an ectatic disorder characterized by progressive corneal thinning and protrusion of its central and/or inferior portions.





What's the diagnosis?

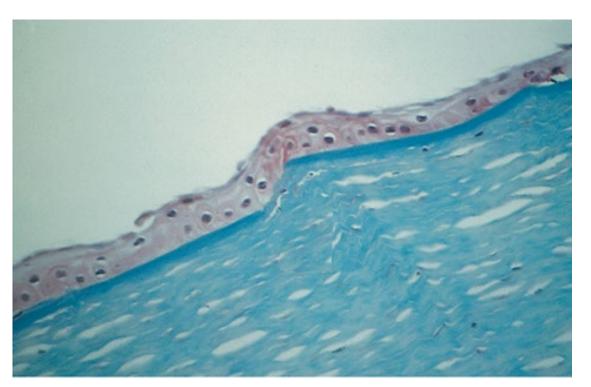
What's going on here? Again saving the obvious, let's note what looks OK:
--The stroma contains the expected artifactual clefts, and isn't picking up stain in a way that suggests it contains some sort of abnormal material.
--While the epithelium looks wonky, it doesn't contain discernible BM running through it.

Now let's talk about the obvious: Bowman's layer is completely disrupted right **here**.

Focal disruption of Bowman's layer in the absence of abnormal stromal staining should bring to mind one dx:

Keratoconus is an ectatic disorder characterized by progressive corneal thinning and protrusion of its central and/or inferior portions. This leads to the characteristic shape (two-words) cornea.





What's the diagnosis?

What's going on here? Again saving the obvious, let's note what looks OK:

--The stroma contains the expected artifactual clefts, and isn't picking up stain in a way that suggests it contains some sort of abnormal material.

--While the epithelium looks wonky, it doesn't contain discernible BM running through it.

Now let's talk about the obvious: Bowman's layer is completely disrupted right **here**.

Focal disruption of Bowman's layer in the absence of abnormal stromal staining should bring to mind one dx:

Keratoconus is an ectatic disorder characterized by progressive corneal thinning and protrusion of its central and/or inferior portions. This leads to the characteristic cone-shaped cornea.





What's going on here? Again saving the obvious, let's note what looks OK:
--The stroma contains the expected artifactual clefts, and isn't picking up stain in a way that suggests it contains some sort of abnormal material.

-While the enithelium looks wonky, it

nning

For more on KCN, see slide-set K38



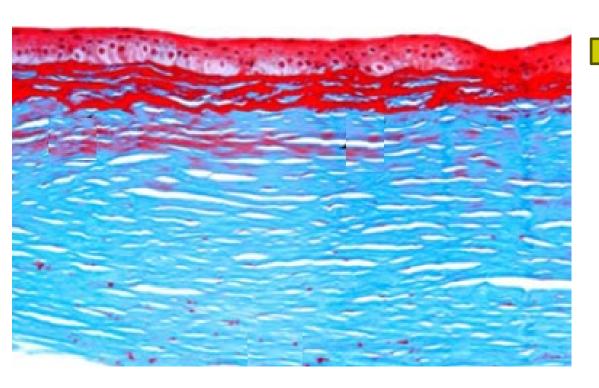
What's the diagnosis?

Now let's talk about the obvious: Bowman's layer is completely disrupted right **here**.

Focal disruption of Bowman's layer in the absence of abnormal stromal staining should bring to mind one dx:

Keratoconus is an ectatic disorder characterized by progressive corneal thinning and protrusion of its central and/or inferior portions. This leads to the characteristic cone-shaped cornea.

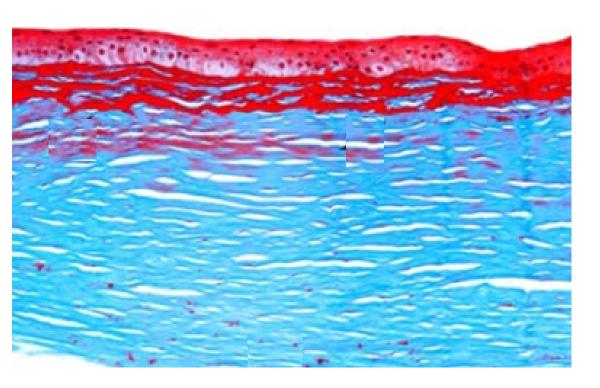




What's going on here? Again, let's first note what looks OK:

--The stroma contains the expected artifactual clefts, so the pathology doesn't involve significant corneal or .

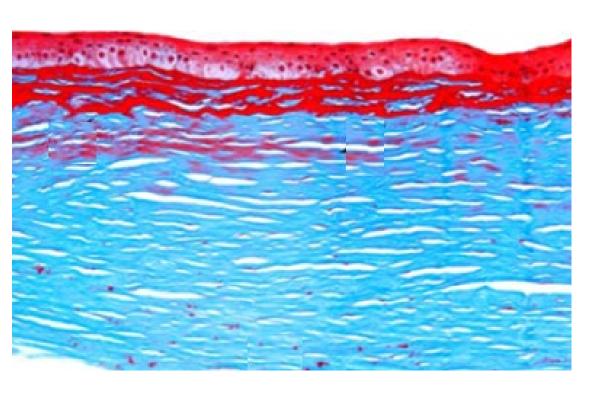




What's going on here? Again, let's first note what looks OK:

--The stroma contains the expected artifactual clefts, so the pathology doesn't involve significant corneal edema or scarring.



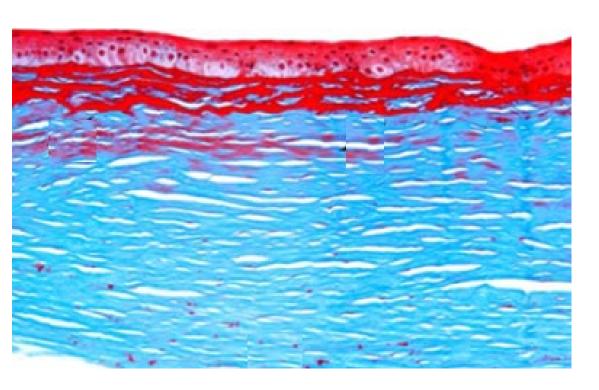


What's going on here? Again, let's first note what looks OK:

--The stroma contains the expected artifactual clefts, so the pathology doesn't involve significant corneal edema or scarring.

That's where the 'normal' ends. Note:
--The epithelium looks wonky, but it
doesn't contain abb. running through it.



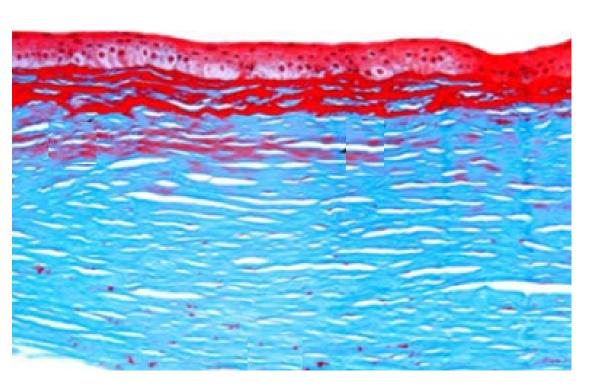


What's going on here? Again, let's first note what looks OK:

--The stroma contains the expected artifactual clefts, so the pathology doesn't involve significant corneal edema or scarring.

That's where the 'normal' ends. Note:
--The epithelium looks wonky, but it
doesn't contain BM running through it.





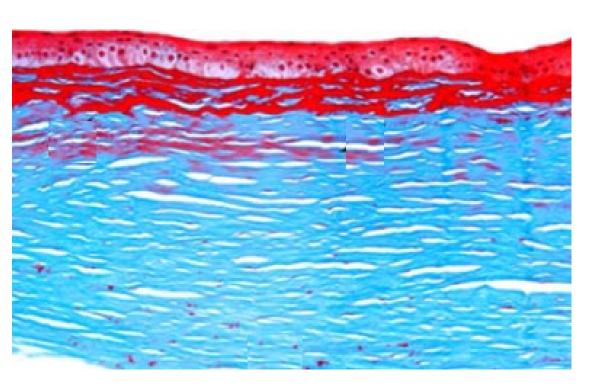
What's going on here? Again, let's first note what looks OK:

--The stroma contains the expected artifactual clefts, so the pathology doesn't involve significant corneal edema or scarring.

That's where the 'normal' ends. Note:

- --The epithelium looks wonky, but it doesn't contain BM running through it.
- -- two words is either gone or severely disrupted





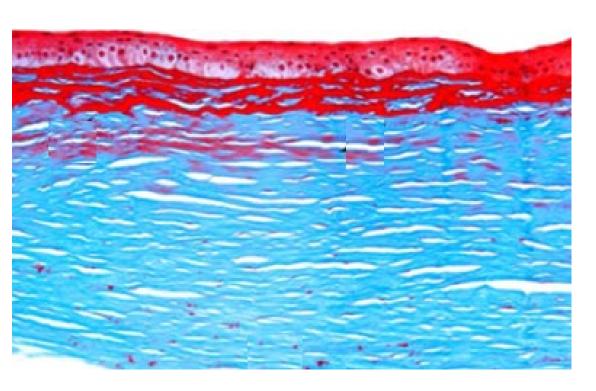
What's going on here? Again, let's first note what looks OK:

--The stroma contains the expected artifactual clefts, so the pathology doesn't involve significant corneal edema or scarring.

That's where the 'normal' ends. Note:

- --The epithelium looks wonky, but it doesn't contain BM running through it.
- --Bowman's layer is either gone or severely disrupted.





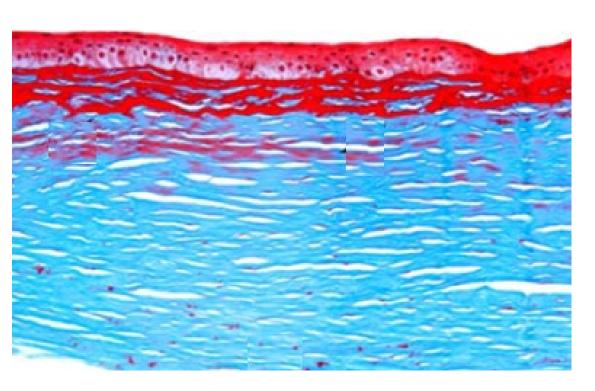
What's going on here? Again, let's first note what looks OK:

--The stroma contains the expected artifactual clefts, so the pathology doesn't involve significant corneal edema or scarring.

That's where the 'normal' ends. Note:

- --The epithelium looks wonky, but it doesn't contain BM running through it.
- --Bowman's layer is either gone or severely disrupted.
- --The stroma is taking a special stain, with the portion that is Bowman's-adjacent staining most avidly.





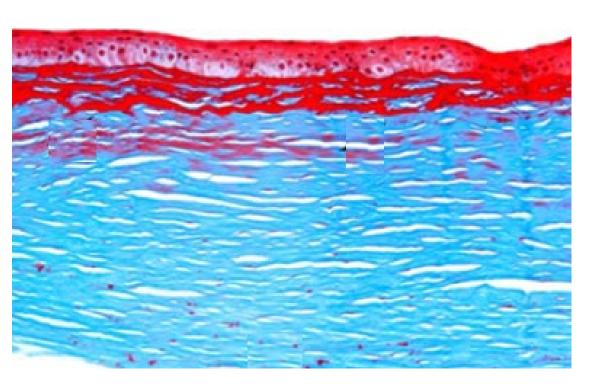
What's going on here? Again, let's first note what looks OK:

--The stroma contains the expected artifactual clefts, so the pathology doesn't involve significant corneal edema or scarring.

That's where the 'normal' ends. Note:

- --The epithelium looks wonky, but it doesn't contain BM running through it.
- --Bowman's layer is either gone or severely disrupted.
- --The stroma is taking a special stain, with the portion that is Bowman's-adjacent staining most avidly. The anterior staining pattern is two-words





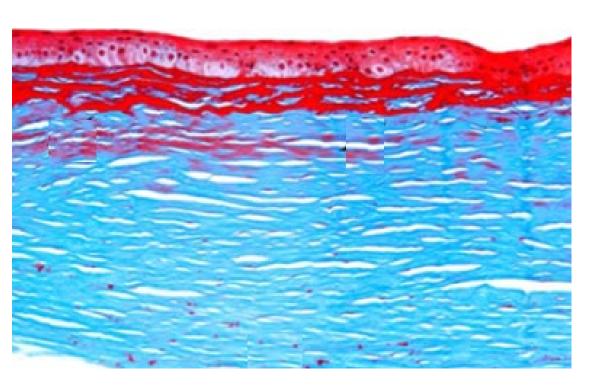
What's going on here? Again, let's first note what looks OK:

--The stroma contains the expected artifactual clefts, so the pathology doesn't involve significant corneal edema or scarring.

That's where the 'normal' ends. Note:

- --The epithelium looks wonky, but it doesn't contain BM running through it.
- --Bowman's layer is either gone or severely disrupted.
- --The stroma is taking a special stain, with the portion that is Bowman's-adjacent staining most avidly. The anterior staining pattern is 'sheet-like.'





What's the diagnosis?

What's going on here? Again, let's first note what looks OK:

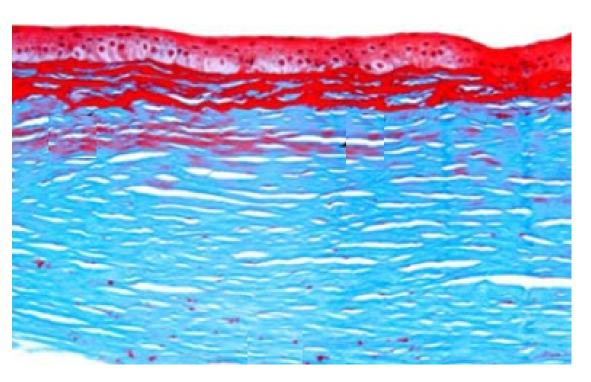
--The stroma contains the expected artifactual clefts, so the pathology doesn't involve significant corneal edema or scarring.

That's where the 'normal' ends. Note:
--The epithelium looks wonky, but it
doesn't contain BM running through it.
--Bowman's layer is either gone or
severely disrupted.

--The stroma is taking a special stain, with the portion that is Bowman's-adjacent staining most avidly. The anterior staining pattern is 'sheet-like.'

Extensive disruption/replacement of Bowman's layer associated with subjacent sheets of avidly-stained material points to one diagnosis:





What's the diagnosis?

What's going on here? Again, let's first note what looks OK:

--The stroma contains the expected artifactual clefts, so the pathology doesn't involve significant corneal edema or scarring.

That's where the 'normal' ends. Note:

- --The epithelium looks wonky, but it doesn't contain BM running through it.
- --Bowman's layer is either gone or severely disrupted.
- --The stroma is taking a special stain, with the portion that is Bowman's-adjacent staining most avidly. The anterior staining pattern is 'sheet-like.'

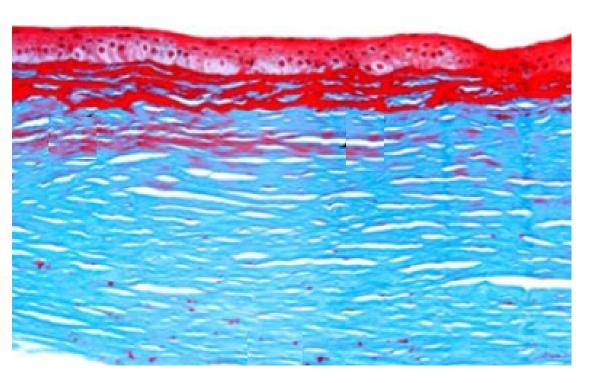
Extensive disruption/replacement of Bowman's layer associated with subjacent sheets of avidly-stained material points to one diagnosis:

Reis-Bücklers corneal dystrophy (RBCD) is one of the

two-words + a long abb.

corneal dystrophies.





What's the diagnosis?

What's going on here? Again, let's first note what looks OK:

--The stroma contains the expected artifactual clefts, so the pathology doesn't involve significant corneal edema or scarring.

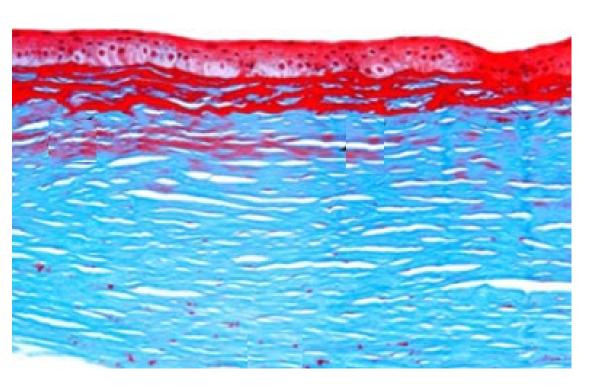
That's where the 'normal' ends. Note:

- --The epithelium looks wonky, but it doesn't contain BM running through it.
- --Bowman's layer is either gone or severely disrupted.
- --The stroma is taking a special stain, with the portion that is Bowman's-adjacent staining most avidly. The anterior staining pattern is 'sheet-like.'

Extensive disruption/replacement of Bowman's layer associated with subjacent sheets of avidly-stained material points to one diagnosis:

Reis-Bücklers corneal dystrophy (RBCD) is one of the epithelial-stromal TGFBI corneal dystrophies.





What's going on here? Again, let's first note what looks OK:

--The stroma contains the expected artifactual clefts, so the pathology doesn't involve significant corneal edema or scarring.

That's where the 'normal' ends. Note:

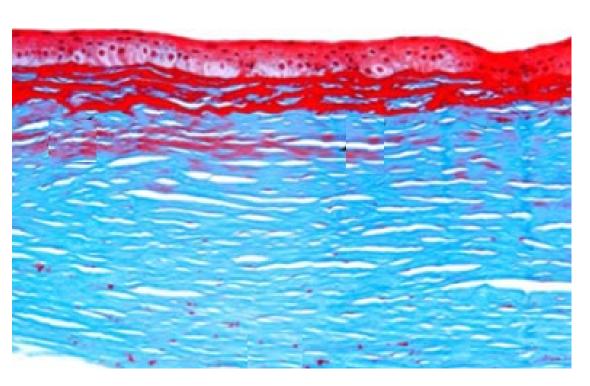
- --The epithelium looks wonky, but it doesn't contain BM running through it.
- --Bowman's layer is either gone or severely disrupted.
- --The stroma is taking a special stain, with the portion that is Bowman's-adjacent staining most avidly. The anterior staining pattern is 'sheet-like.'

Extensive disruption/replacement of Bowman's layer associated with subjacent sheets of avidly-stained material points to one diagnosis:

What's the diagnosis?

Reis-Bücklers corneal dystrophy (RBCD) is one of the *epithelial-stromal TGFBI* corneal dystrophies. It mainly affects two words





What's going on here? Again, let's first note what looks OK:

--The stroma contains the expected artifactual clefts, so the pathology doesn't involve significant corneal edema or scarring.

That's where the 'normal' ends. Note:

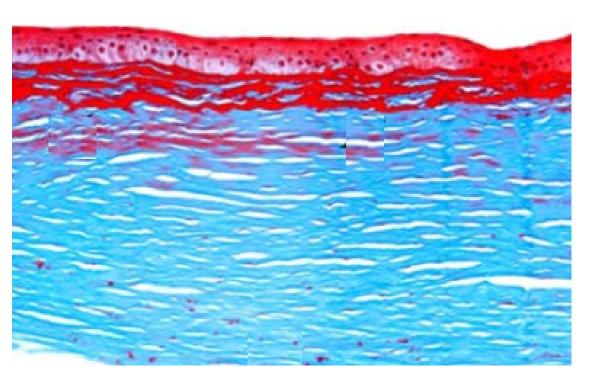
- --The epithelium looks wonky, but it doesn't contain BM running through it.
- --Bowman's layer is either gone or severely disrupted.
- --The stroma is taking a special stain, with the portion that is Bowman's-adjacent staining most avidly. The anterior staining pattern is 'sheet-like.'

Extensive disruption/replacement of Bowman's layer associated with subjacent sheets of avidly-stained material points to one diagnosis:

What's the diagnosis?

Reis-Bücklers corneal dystrophy (RBCD) is one of the *epithelial-stromal TGFBI* corneal dystrophies. It mainly affects Bowman's layer*.





What's going on here? Again, let's first note what looks OK:

--The stroma contains the expected artifactual clefts, so the pathology doesn't involve significant corneal edema or scarring.

That's where the 'normal' ends. Note:

- --The epithelium looks wonky, but it doesn't contain BM running through it.
- --Bowman's layer is either gone or severely disrupted.
- --The stroma is taking a special stain, with the portion that is Bowman's-adjacent staining most avidly. The anterior staining pattern is 'sheet-like.'

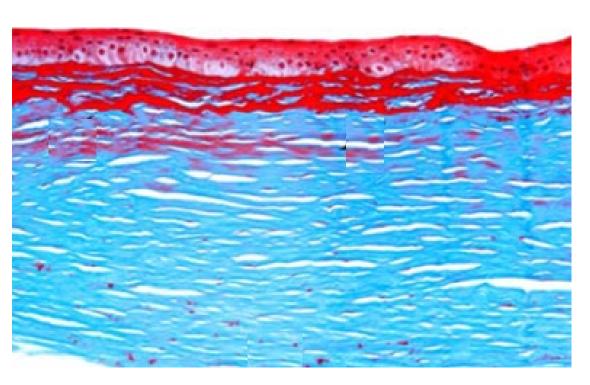
Extensive disruption/replacement of Bowman's layer associated with subjacent sheets of avidly-stained material points to one diagnosis:

What's the diagnosis?

Reis-Bücklers corneal dystrophy (RBCD) is one of the *epithelial-stromal TGFBI* corneal dystrophies. It mainly affects Bowman's layer*.

*Under the classification system used in previous volumes of the *Cornea* book, RBCD was classified as a *Corneal Dystrophy of Bowman's* (CDB).





What's the diagnosis?

What's going on here? Again, let's first note what looks OK:

--The stroma contains the expected artifactual clefts, so the pathology doesn't involve significant corneal edema or scarring.

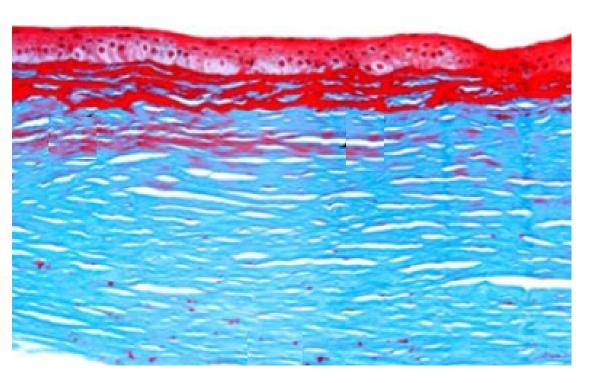
That's where the 'normal' ends. Note:

- --The epithelium looks wonky, but it doesn't contain BM running through it.
- --Bowman's layer is either gone or severely disrupted.
- --The stroma is taking a special stain, with the portion that is Bowman's-adjacent staining most avidly. The anterior staining pattern is 'sheet-like.'

Extensive disruption/replacement of Bowman's layer associated with subjacent sheets of avidly-stained material points to one diagnosis:

Reis-Bücklers corneal dystrophy (RBCD) is one of the *epithelial-stromal TGFBI* corneal dystrophies. It mainly affects Bowman's layer* . Primary complaints are related to three words.





What's the diagnosis?

What's going on here? Again, let's first note what looks OK:

--The stroma contains the expected artifactual clefts, so the pathology doesn't involve significant corneal edema or scarring.

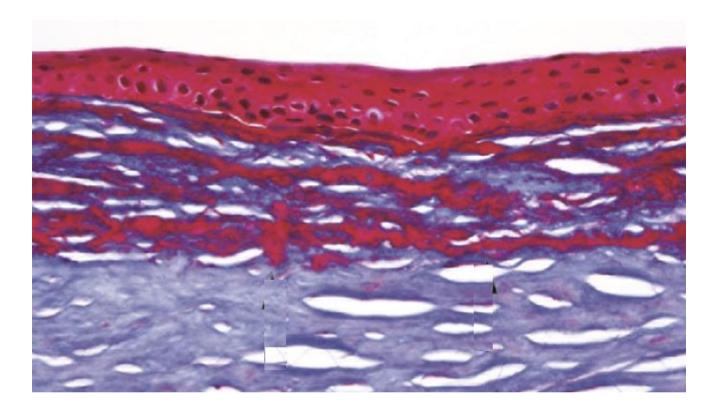
That's where the 'normal' ends. Note:

- --The epithelium looks wonky, but it doesn't contain BM running through it.
- --Bowman's layer is either gone or severely disrupted.
- --The stroma is taking a special stain, with the portion that is Bowman's-adjacent staining most avidly. The anterior staining pattern is 'sheet-like.'

Extensive disruption/replacement of Bowman's layer associated with subjacent sheets of avidly-stained material points to one diagnosis:

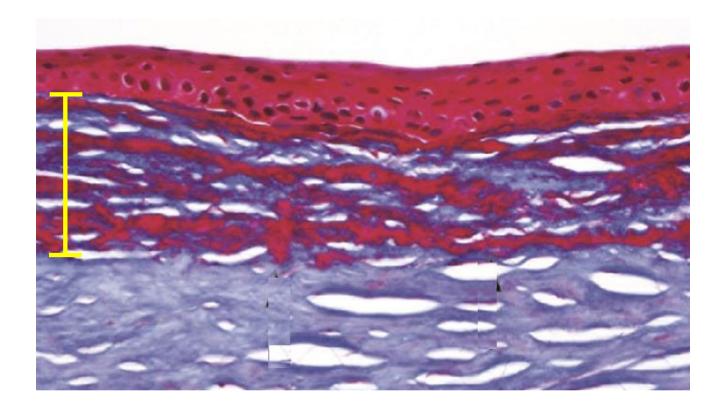
Reis-Bücklers corneal dystrophy (RBCD) is one of the *epithelial-stromal TGFBI* corneal dystrophies. It mainly affects Bowman's layer* . Primary complaints are related to recurrent epithelial erosions .



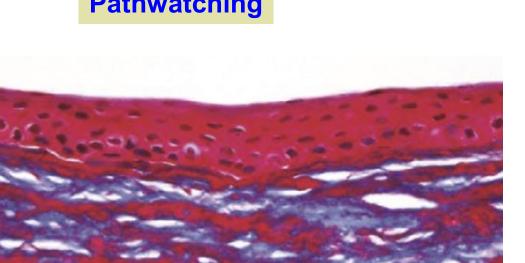


Another characteristic of RBCD: The inverse relationship between stromal involvement and the robustness of the overlying epithelium.





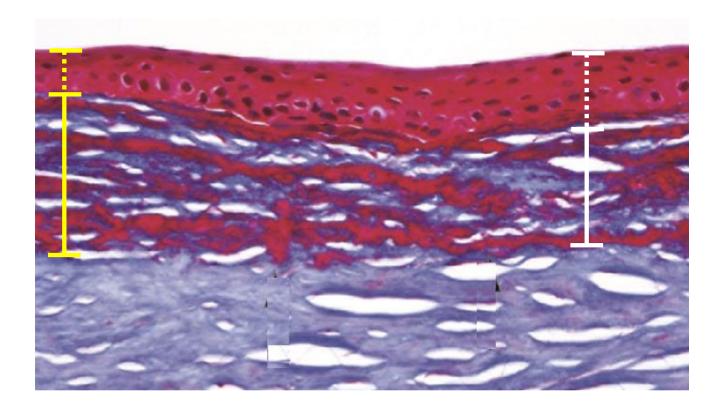
Another characteristic of RBCD: The inverse relationship between stromal involvement and the robustness of the overlying epithelium. That is, note that where the stromal involvement is greater...





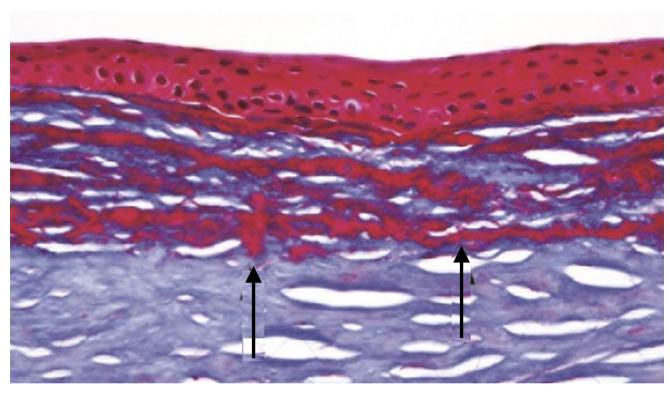
Another characteristic of RBCD: The inverse relationship between stromal involvement and the robustness of the overlying epithelium. That is, note that where the stromal involvement is greater...the overlying epithelium is *thinner*.





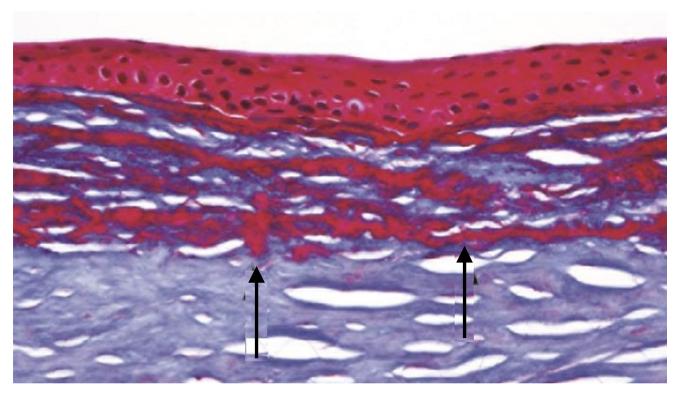
Another characteristic of RBCD: The inverse relationship between stromal involvement and the robustness of the overlying epithelium. That is, note that where the stromal involvement is greater...the overlying epithelium is *thinner*. The reverse is also true.





Note: This image appears in both the *Cornea* and *Path* books. Here is how it is captioned in each:

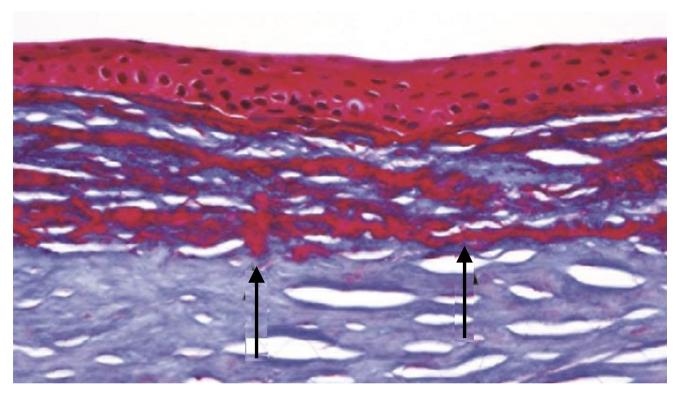




Note: This image appears in both the *Cornea* and *Path* books. Here is how it is captioned in each:

<u>In the Cornea book</u>: Light microscopy with Masson trichrome stain reveals replacement of Bowman's layer (arrows) with hyaline. Note thinner epithelium overlying areas of increased stromal involvement and vice-versa



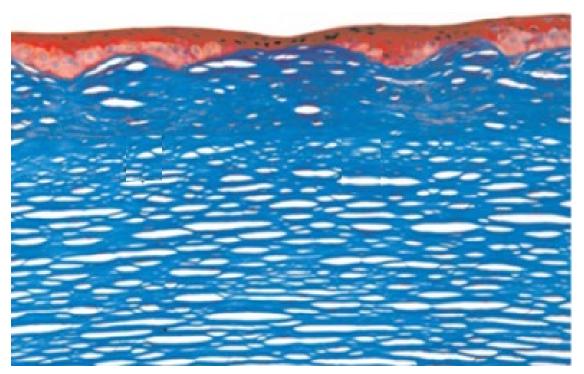


Note: This image appears in both the *Cornea* and *Path* books. Here is how it is captioned in each:

<u>In the Cornea book</u>: Light microscopy with Masson trichrome stain reveals replacement of Bowman's layer (arrows) with hyaline. Note thinner epithelium overlying areas of increased stromal involvement and vice-versa

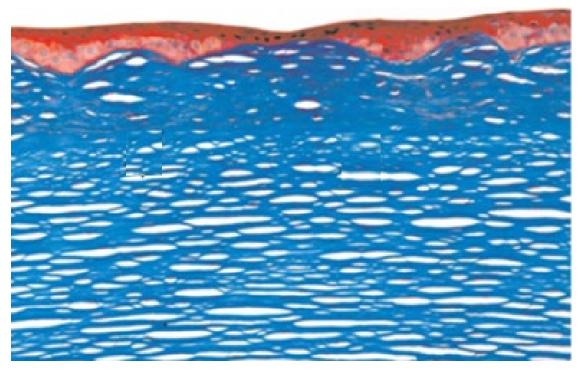
<u>In the Path book</u>: Masson trichrome stain demonstrates diffuse loss of Bowman layer, superficial stromal fibrosis, and numerous red deposits (arrows).





There is one more epithelial-stromal TGFBI dystrophy that primarily effects Bowman's (pictured)—what is it?

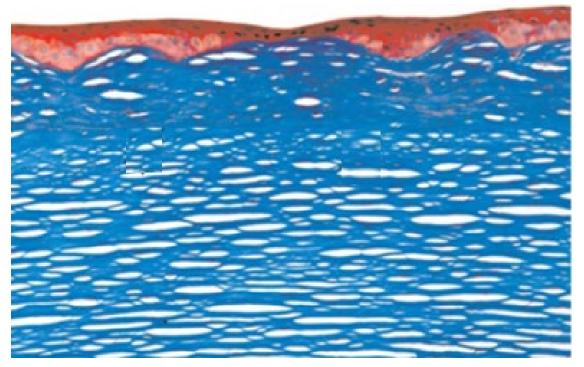




Thiel-Behnke corneal dystrophy (TBCD)

There is one more epithelial-stromal TGFBI dystrophy that primarily effects Bowman's (pictured)—what is it?
Thiel-Behnke corneal dystrophy (TBCD)



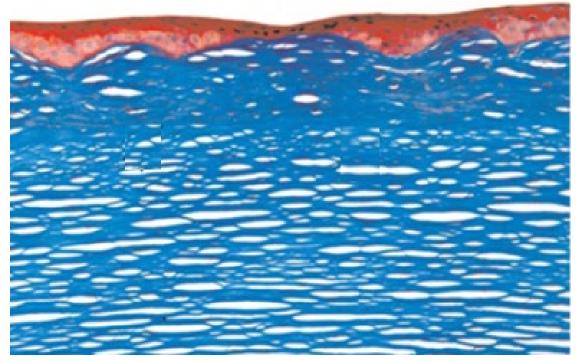


Thiel-Behnke corneal dystrophy (TBCD)

There is one more epithelial-stromal TGFBI dystrophy that primarily effects Bowman's (pictured)—what is it?
Thiel-Behnke corneal dystrophy (TBCD)

Was TBCD previously a CDB a la RBCD?



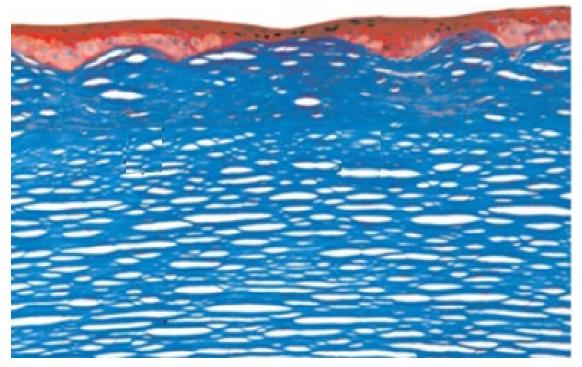


Thiel-Behnke corneal dystrophy (TBCD)

There is one more epithelial-stromal TGFBI dystrophy that primarily effects Bowman's (pictured)—what is it?
Thiel-Behnke corneal dystrophy (TBCD)

Was TBCD previously a CDB a la RBCD? Indeed it was





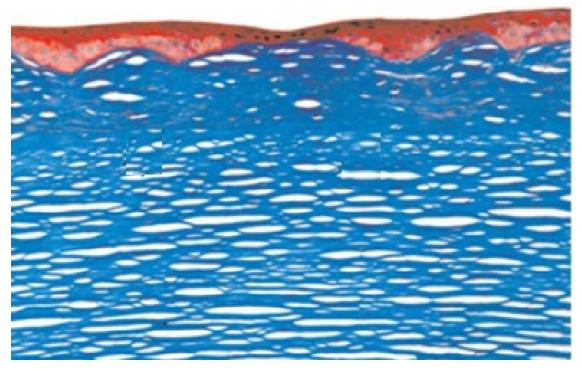
Thiel-Behnke corneal dystrophy (TBCD)

There is one more epithelial-stromal TGFBI dystrophy that primarily effects Bowman's (pictured)—what is it?
Thiel-Behnke corneal dystrophy (TBCD)

Was TBCD previously a CDB a la RBCD? Indeed it was

Does TBCD replace Bowman's with abnormal material a la RBCD?





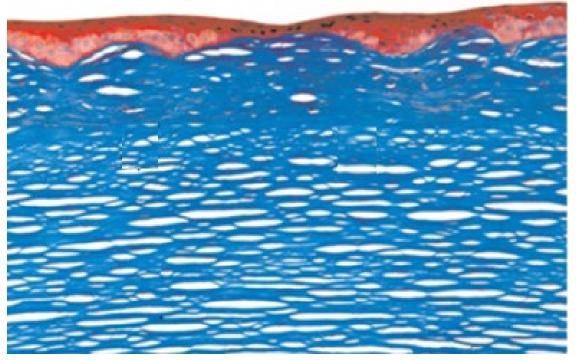
Thiel-Behnke corneal dystrophy (TBCD)

There is one more epithelial-stromal TGFBI dystrophy that primarily effects Bowman's (pictured)—what is it?
Thiel-Behnke corneal dystrophy (TBCD)

Was TBCD previously a CDB a la RBCD? Indeed it was

Does TBCD replace Bowman's with abnormal material a la RBCD? Indeed it does





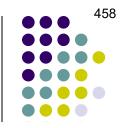
Thiel-Behnke corneal dystrophy (TBCD)

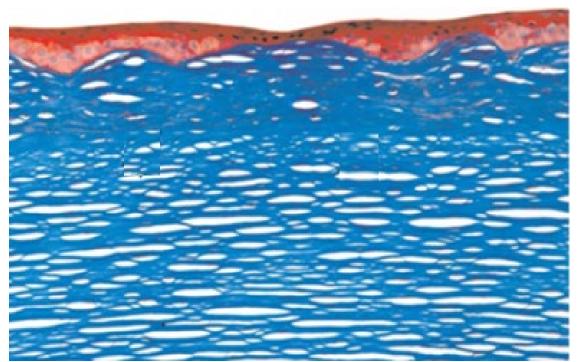
There is one more epithelial-stromal TGFBI dystrophy that primarily effects Bowman's (pictured)—what is it?
Thiel-Behnke corneal dystrophy (TBCD)

Was TBCD previously a CDB a la RBCD? Indeed it was

Does TBCD replace Bowman's with abnormal material a la RBCD? Indeed it does

With respect to their path features, how can they be differentiated?





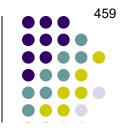
Thiel-Behnke corneal dystrophy (TBCD)

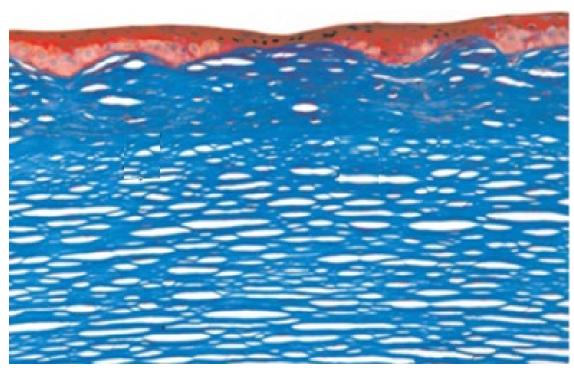
There is one more epithelial-stromal TGFBI dystrophy that primarily effects Bowman's (pictured)—what is it?
Thiel-Behnke corneal dystrophy (TBCD)

Was TBCD previously a CDB a la RBCD? Indeed it was

Does TBCD replace Bowman's with abnormal material a la RBCD? Indeed it does

With respect to their path features, how can they be differentiated? In photomicrographs: By the form taken by the abnormal material.





Thiel-Behnke corneal dystrophy (TBCD)

There is one more epithelial-stromal TGFBI dystrophy that primarily effects Bowman's (pictured)—what is it?

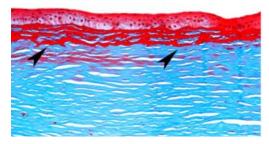
Thiel-Behnke corneal dystrophy (TBCD)

Was TBCD previously a CDB a la RBCD? Indeed it was

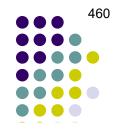
Does TBCD replace Bowman's with abnormal material a la RBCD? Indeed it does

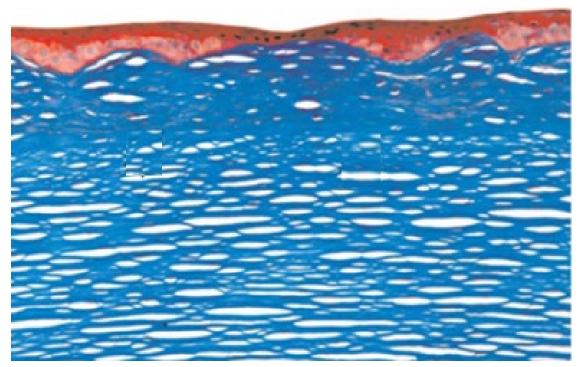
With respect to their path features, how can they be differentiated?

In <u>photomicrographs</u>: By the form taken by the abnormal material. Recall that in RBCD the material was described as ...



RBCD





Thiel-Behnke corneal dystrophy (TBCD)

There is one more epithelial-stromal TGFBI dystrophy that primarily effects Bowman's (pictured)—what is it?

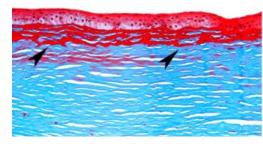
Thiel-Behnke corneal dystrophy (TBCD)

Was TBCD previously a CDB a la RBCD? Indeed it was

Does TBCD replace Bowman's with abnormal material a la RBCD? Indeed it does

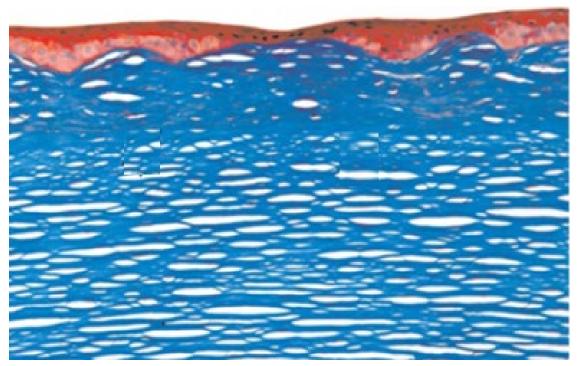
With respect to their path features, how can they be differentiated?

In <u>photomicrographs</u>: By the form taken by the abnormal material. Recall that in RBCD the material was described as 'layered'.



RBCD





Thiel-Behnke corneal dystrophy (TBCD)

There is one more epithelial-stromal TGFBI dystrophy that primarily effects Bowman's (pictured)—what is it?

Thiel-Behnke corneal dystrophy (TBCD)

Was TBCD previously a CDB a la RBCD? Indeed it was

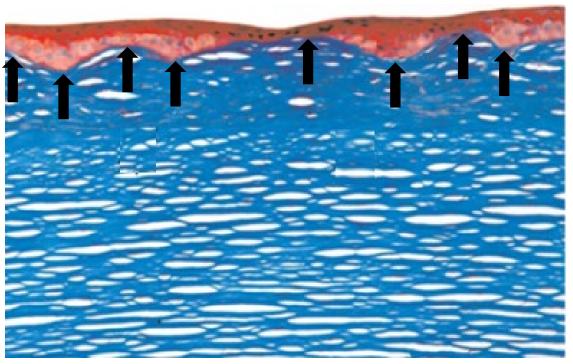
Does TBCD replace Bowman's with abnormal material a la RBCD? Indeed it does

With respect to their path features, how can they be differentiated?

In photomicrographs: By the form taken by the abnormal material. Recall that in RBCD the material was described as 'layered'.

Contrast that with TBCD in which the form is described as buzzword





Thiel-Behnke corneal dystrophy (TBCD)

There is one more epithelial-stromal TGFBI dystrophy that primarily effects Bowman's (pictured)—what is it?

Thiel-Behnke corneal dystrophy (TBCD)

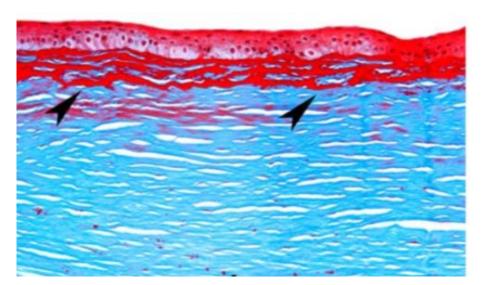
Was TBCD previously a CDB a la RBCD? Indeed it was

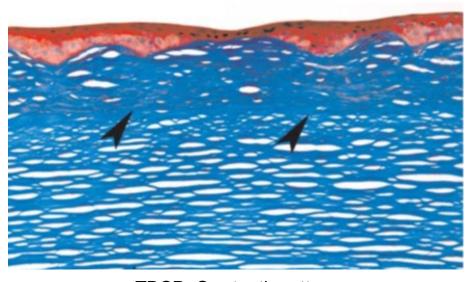
Does TBCD replace Bowman's with abnormal material a la RBCD? Indeed it does

With respect to their path features, how can they be differentiated?

In <u>photomicrographs</u>: By the form taken by the abnormal material. Recall that in RBCD the material was described as 'layered'. Contrast that with TBCD in which the form is described as 'sawtooth'.





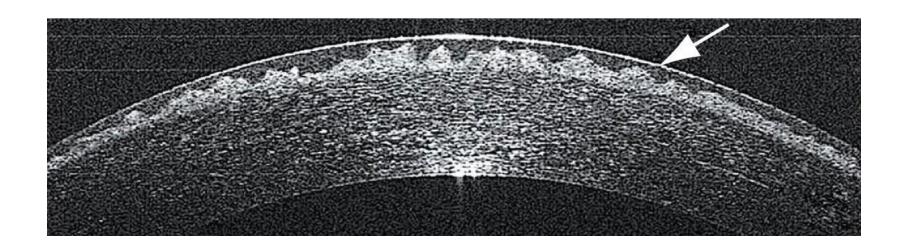


RBCD: sheet-like layers

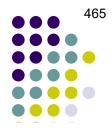
TBCD: Sawtooth pattern

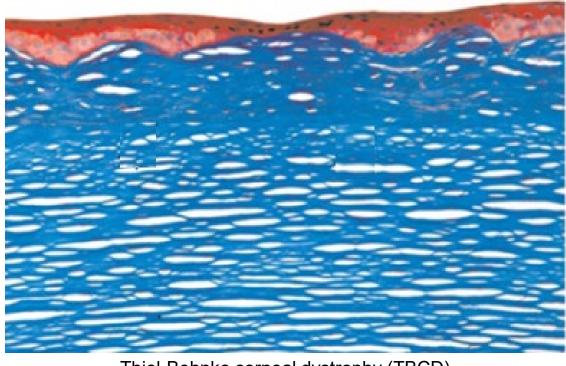
RBCD and TBCD: Photomicrographs demonstrating their characteristic forms





TBCD. The sawtooth pattern is readily apparent on anterior-segment OCT.





Thiel-Behnke corneal dystrophy (TBCD)

Next Q

There is one more epithelial-stromal TGFBI dystrophy that primarily effects Bowman's (pictured)—what is it?
Thiel-Behnke corneal dystrophy (TBCD)

Was TBCD previously a CDB a la RBCD? Indeed it was

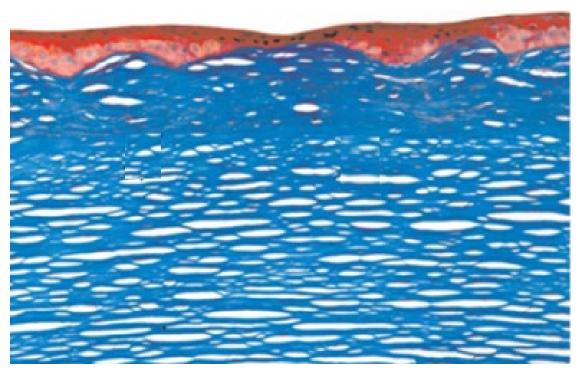
Does TBCD replace Bowman's with abnormal material a la RBCD? Indeed it does

With respect to their path features, how can they be differentiated?

In <u>photomicrographs</u>: By the form taken by the abnormal material. Recall that in RBCD the material was described as 'layered'. Contrast that with TBCD in which the form is described as 'sawtooth'.

In <u>electron microscopy</u>: By the <u>property</u> of the fibers comprising the abnormal material





Thiel-Behnke corneal dystrophy (TBCD)

There is one more epithelial-stromal TGFBI dystrophy that primarily effects Bowman's (pictured)—what is it?

Thiel-Behnke corneal dystrophy (TBCD)

Was TBCD previously a CDB a la RBCD? Indeed it was

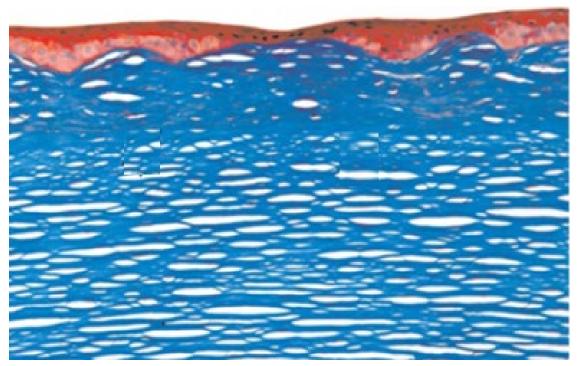
Does TBCD replace Bowman's with abnormal material a la RBCD? Indeed it does

With respect to their path features, how can they be differentiated?

In <u>photomicrographs</u>: By the form taken by the abnormal material. Recall that in RBCD the material was described as 'layered'. Contrast that with TBCD in which the form is described as 'sawtooth'.

In <u>electron microscopy</u>: By the shape of the fibers comprising the abnormal material





Thiel-Behnke corneal dystrophy (TBCD)

There is one more epithelial-stromal TGFBI dystrophy that primarily effects Bowman's (pictured)—what is it?

Thiel-Behnke corneal dystrophy (TBCD)

Was TBCD previously a CDB a la RBCD? Indeed it was

Does TBCD replace Bowman's with abnormal material a la RBCD? Indeed it does

With respect to their path features, how can they be differentiated?

In <u>photomicrographs</u>: By the form taken by the abnormal material. Recall that in RBCD the material was described as 'layered'. Contrast that with TBCD in which the form is described as 'sawtooth'.

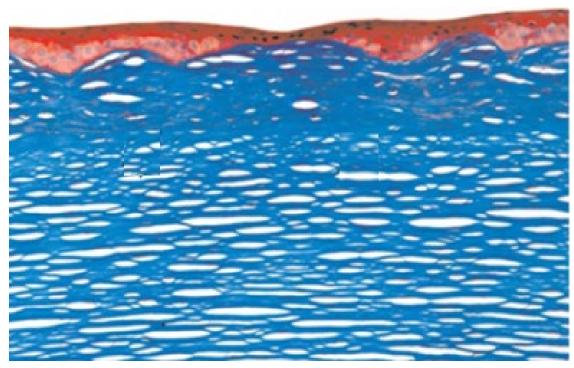
In <u>electron microscopy</u>: By the shape of the fibers comprising the abnormal material:

--In RCBD, fibers are

two-words

--In TBCD fibers are





Thiel-Behnke corneal dystrophy (TBCD)

There is one more epithelial-stromal TGFBI dystrophy that primarily effects Bowman's (pictured)—what is it?

Thiel-Behnke corneal dystrophy (TBCD)

Was TBCD previously a CDB a la RBCD? Indeed it was

Does TBCD replace Bowman's with abnormal material a la RBCD? Indeed it does

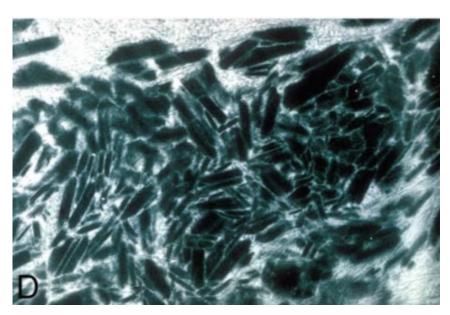
With respect to their path features, how can they be differentiated?

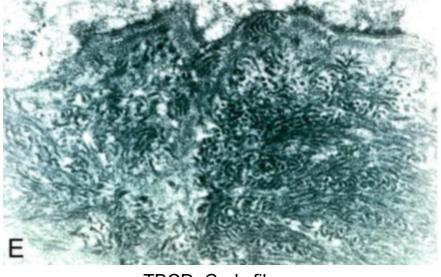
In <u>photomicrographs</u>: By the form taken by the abnormal material. Recall that in RBCD the material was described as 'layered'. Contrast that with TBCD in which the form is described as 'sawtooth'.

In <u>electron microscopy</u>: By the shape of the fibers comprising the abnormal material:

- --In RCBD, fibers are rod-shaped
- --In TBCD fibers are curly



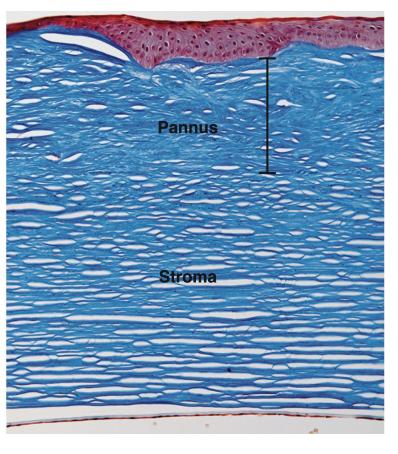




RBCD: Rod-shaped fibers

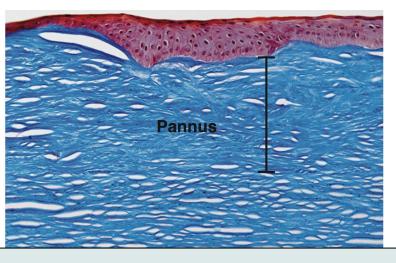
TBCD: Curly fibers

RBCD and TBCD: Electron microscopy



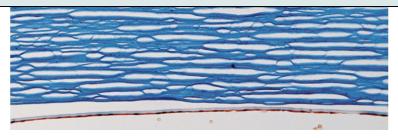


<u>From the Path* book</u>: Thiel-Behnke corneal dystrophy. Masson trichrome stain demonstrates diffuse replacement of Bowman layer by a thick fibrous pannus (*bracket*). The overlying epithelium exhibits a sawtooth configuration. The underlying stroma appears to be uninvolved.





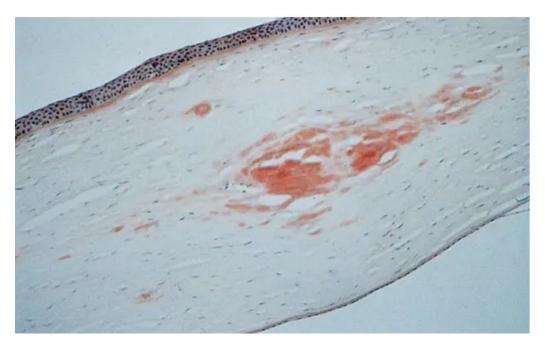
For more on RBCD and TBCD, see slide-set K42



<u>From the Path* book</u>: Thiel-Behnke corneal dystrophy. Masson trichrome stain demonstrates diffuse replacement of Bowman layer by a thick fibrous pannus *(bracket)*. The overlying epithelium exhibits a sawtooth configuration. The underlying stroma appears to be uninvolved.

*(The most recent version of the *Cornea* book does not contain a TBCD photomicrograph)

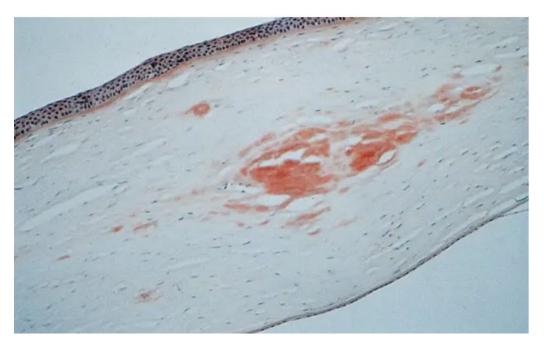




What's going on here? Again, let's first note what looks OK:

--The epithelium seems to be unaffected --It's hard to see Bowman's the whole way across, and small sections may be hinky. But for the most part it appears intact—certainly lacking evidence of the extensive disruption we'd expect from a CDB or keratoconus.

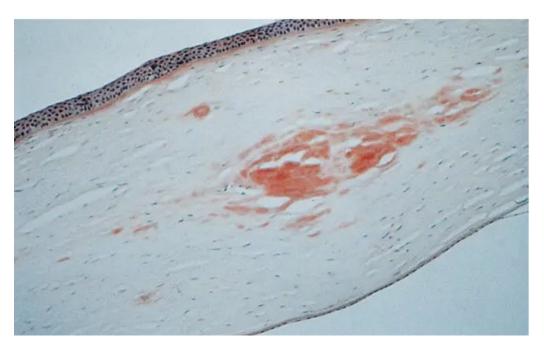




What's going on here? Again, let's first note what looks OK:

- --The epithelium seems to be unaffected --It's hard to see Bowman's the whole way across, and small sections may be hinky. But for the most part it appears intact—certainly lacking evidence of the extensive disruption we'd expect from a CDB or keratoconus. Now the wonky:
- --Stromal deposits, mainly in the central and anterior regions, are taking a stain.

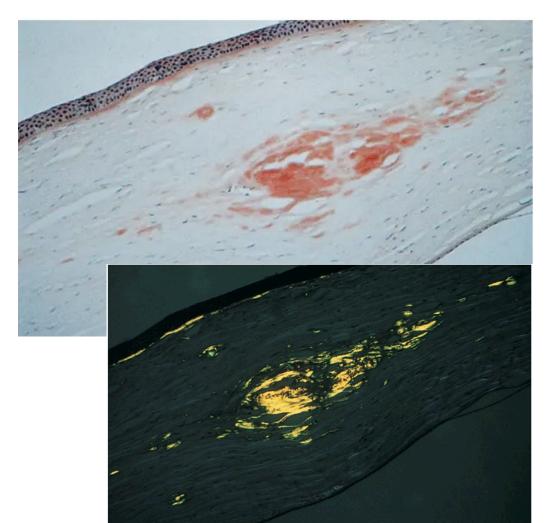




What's going on here? Again, let's first note what looks OK:

- --The epithelium seems to be unaffected --It's hard to see Bowman's the whole way across, and small sections may be hinky. But for the most part it appears intact—certainly lacking evidence of the extensive disruption we'd expect from a CDB or keratoconus. Now the wonky:
- --Stromal deposits, mainly in the central and anterior regions, are taking a stain.
- --The expected stromal clefts are missing around a glom of material, indicating distortion of the adjacent cornea's lamellar structure.





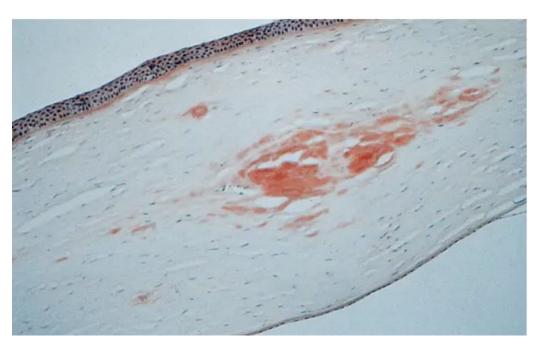
What's going on here? Again, let's first note what looks OK:

- --The epithelium seems to be unaffected --It's hard to see Bowman's the whole way across, and small sections may be hinky. But for the most part it appears intact—certainly lacking evidence of the extensive disruption we'd expect from a CDB or keratoconus. Now the wonky:
- --Stromal deposits, mainly in the central and anterior regions, are taking a stain.
- --The expected stromal clefts are missing around a glom of material, indicating distortion of the adjacent cornea's lamellar structure.

 This may be presented concurrently—the

same slice of cornea under a different illumination in which the material glows green.





What's the diagnosis?

What's going on here? Again, let's first note what looks OK:

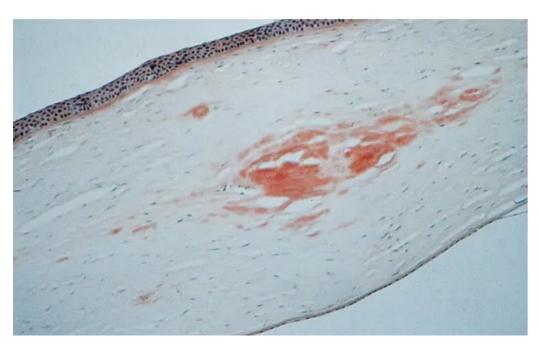
- --The epithelium seems to be unaffected --It's hard to see Bowman's the whole way across, and small sections may be hinky. But for the most part it appears intact—certainly lacking evidence of the extensive disruption we'd expect from a CDB or keratoconus. Now the wonky:
- --Stromal deposits, mainly in the central and anterior regions, are taking a stain.
- --The expected stromal clefts are missing around a glom of material, indicating distortion of the adjacent cornea's lamellar structure.

 This may be presented concurrently—the same slice of cornea under a different

The presence of anterior/mid-stromal material that stains red but glows green can be only one thing:

illumination in which the material glows green.





What's the diagnosis?

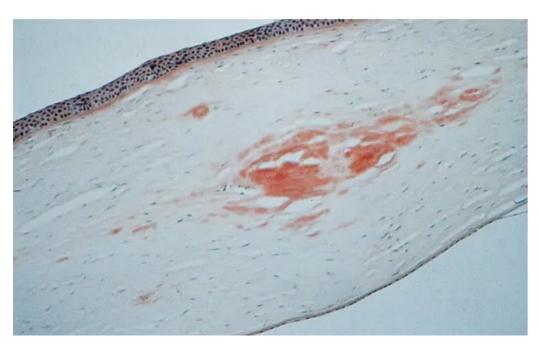
What's going on here? Again, let's first note what looks OK:

- --The epithelium seems to be unaffected --It's hard to see Bowman's the whole way across, and small sections may be hinky. But for the most part it appears intact—certainly lacking evidence of the extensive disruption we'd expect from a CDB or keratoconus. Now the wonky:
- --Stromal deposits, mainly in the central and anterior regions, are taking a stain.
- --The expected stromal clefts are missing around a glom of material, indicating distortion of the adjacent cornea's lamellar structure.

 This may be presented concurrently—the same slice of cornea under a different illumination in which the material glows green. The presence of anterior/mid-stromal material that stains red but glows green can be only one thing:

Lattice corneal dystrophy (LCD) one of the epithelial-stromal TGFBI corneal dystrophies.





What's the diagnosis?

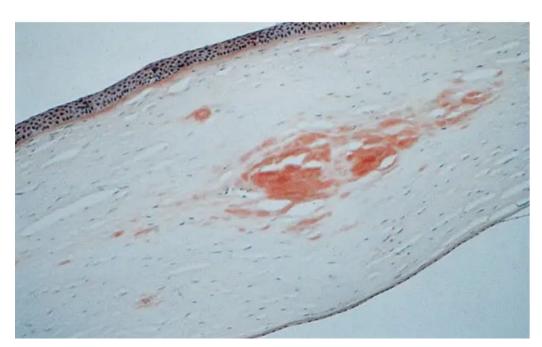
What's going on here? Again, let's first note what looks OK:

- --The epithelium seems to be unaffected --It's hard to see Bowman's the whole way across, and small sections may be hinky. But for the most part it appears intact—certainly lacking evidence of the extensive disruption we'd expect from a CDB or keratoconus. Now the wonky:
- --Stromal deposits, mainly in the central and anterior regions, are taking a stain.
- --The expected stromal clefts are missing around a glom of material, indicating distortion of the adjacent cornea's lamellar structure.

 This may be presented concurrently—the same slice of cornea under a different illumination in which the material glows green. The presence of anterior/mid-stromal material that stains red but glows green can be only one thing:

Lattice corneal dystrophy (LCD) is one of the epithelial-stromal TGFBI corneal dystrophies.





What's the diagnosis?

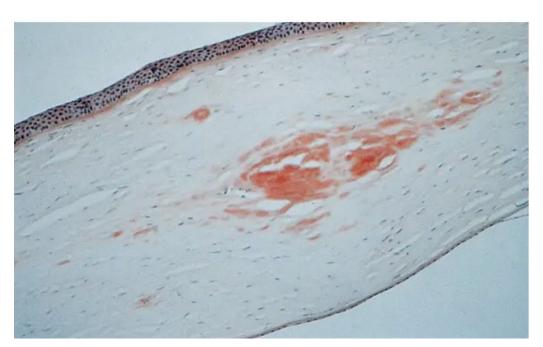
What's going on here? Again, let's first note what looks OK:

- --The epithelium seems to be unaffected --It's hard to see Bowman's the whole way across, and small sections may be hinky. But for the most part it appears intact—certainly lacking evidence of the extensive disruption we'd expect from a CDB or keratoconus. Now the wonky:
- --Stromal deposits, mainly in the central and anterior regions, are taking a stain.
- --The expected stromal clefts are missing around a glom of material, indicating distortion of the adjacent cornea's lamellar structure.

 This may be presented concurrently—the same slice of cornea under a different illumination in which the material glows green. The presence of anterior/mid-stromal material that stains red but glows green can be only one thing:

Lattice corneal dystrophy (LCD) is one of the *epithelial-stromal TGFBI* corneal dystrophies. The *BCSC* recognizes # variants





What's the diagnosis?

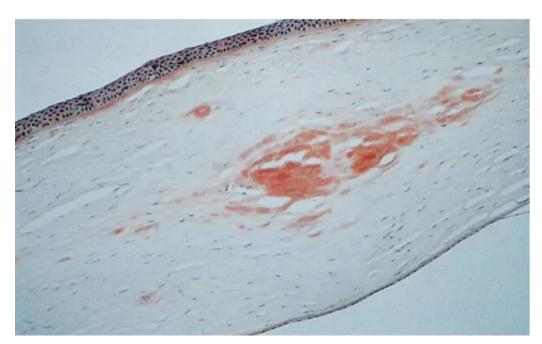
What's going on here? Again, let's first note what looks OK:

- --The epithelium seems to be unaffected --It's hard to see Bowman's the whole way across, and small sections may be hinky. But for the most part it appears intact—certainly lacking evidence of the extensive disruption we'd expect from a CDB or keratoconus. Now the wonky:
- --Stromal deposits, mainly in the central and anterior regions, are taking a stain.
- --The expected stromal clefts are missing around a glom of material, indicating distortion of the adjacent cornea's lamellar structure.

 This may be presented concurrently—the same slice of cornea under a different illumination in which the material glows green. The presence of anterior/mid-stromal material that stains red but glows green can be only one thing:

Lattice corneal dystrophy (LCD) is one of the *epithelial-stromal TGFBI* corneal dystrophies. The *BCSC* recognizes five variants





What's the diagnosis?

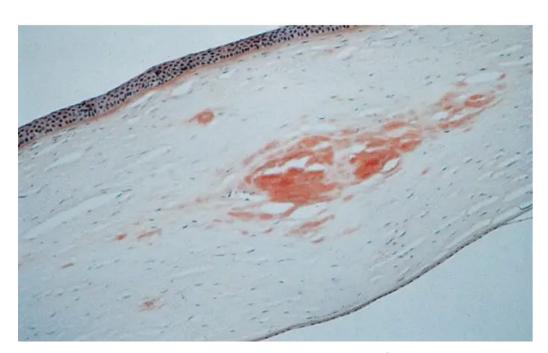
What's going on here? Again, let's first note what looks OK:

- --The epithelium seems to be unaffected --It's hard to see Bowman's the whole way across, and small sections may be hinky. But for the most part it appears intact—certainly lacking evidence of the extensive disruption we'd expect from a CDB or keratoconus. Now the wonky:
- --Stromal deposits, mainly in the central and anterior regions, are taking a stain.
- --The expected stromal clefts are missing around a glom of material, indicating distortion of the adjacent cornea's lamellar structure.

 This may be presented concurrently—the same slice of cornea under a different illumination in which the material glows green. The presence of anterior/mid-stromal material that stains red but glows green can be only one thing:

Lattice corneal dystrophy (LCD) is one of the *epithelial-stromal TGFBI* corneal dystrophies. The *BCSC* recognizes five variants, but spends essentially all of its attention on the fine strong (aka lattice).





What's the diagnosis?

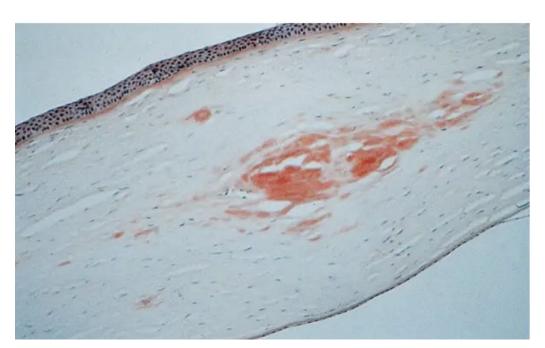
What's going on here? Again, let's first note what looks OK:

- --The epithelium seems to be unaffected --It's hard to see Bowman's the whole way across, and small sections may be hinky. But for the most part it appears intact—certainly lacking evidence of the extensive disruption we'd expect from a CDB or keratoconus. Now the wonky:
- --Stromal deposits, mainly in the central and anterior regions, are taking a stain.
- --The expected stromal clefts are missing around a glom of material, indicating distortion of the adjacent cornea's lamellar structure.

 This may be presented concurrently—the same slice of cornea under a different illumination in which the material glows green. The presence of anterior/mid-stromal material that stains red but glows green can be only one thing:

Lattice corneal dystrophy (LCD) is one of the *epithelial-stromal TGFBI* corneal dystrophies. The *BCSC* recognizes five variants, but spends essentially all of its attention on LCD1 (aka 'classic' lattice).





What's the diagnosis?

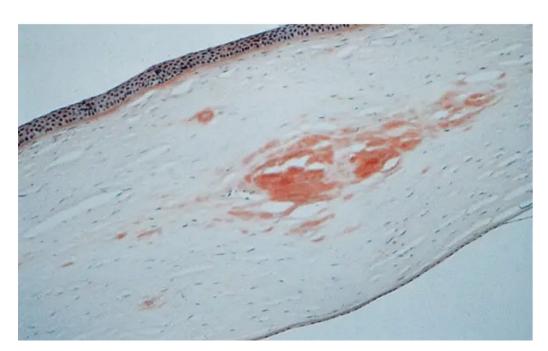
What's going on here? Again, let's first note what looks OK:

- --The epithelium seems to be unaffected --It's hard to see Bowman's the whole way across, and small sections may be hinky. But for the most part it appears intact—certainly lacking evidence of the extensive disruption we'd expect from a CDB or keratoconus. Now the wonky:
- --Stromal deposits, mainly in the central and anterior regions, are taking a stain.
- --The expected stromal clefts are missing around a glom of material, indicating distortion of the adjacent cornea's lamellar structure.

 This may be presented concurrently—the same slice of cornea under a different illumination in which the material glows green. The presence of anterior/mid-stromal material that stains red but glows green can be only one thing:

Lattice corneal dystrophy (LCD) is one of the *epithelial-stromal TGFBI* corneal dystrophies. The *BCSC* recognizes five variants, but spends essentially all of its attention on LCD1 (aka 'classic' lattice). The LCD pathologic process involves the deposition of in the cornea, mainly in the mid- and anterior stroma.





What's the diagnosis?

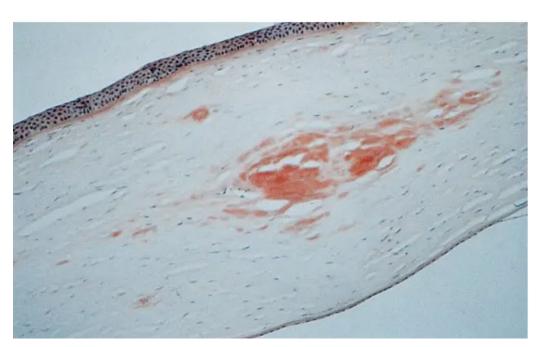
What's going on here? Again, let's first note what looks OK:

- --The epithelium seems to be unaffected --It's hard to see Bowman's the whole way across, and small sections may be hinky. But for the most part it appears intact—certainly lacking evidence of the extensive disruption we'd expect from a CDB or keratoconus. Now the wonky:
- --Stromal deposits, mainly in the central and anterior regions, are taking a stain.
- --The expected stromal clefts are missing around a glom of material, indicating distortion of the adjacent cornea's lamellar structure.

 This may be presented concurrently—the same slice of cornea under a different illumination in which the material glows green. The presence of anterior/mid-stromal material that stains red but glows green can be only one thing:

Lattice corneal dystrophy (LCD) is one of the *epithelial-stromal TGFBI* corneal dystrophies. The *BCSC* recognizes five variants, but spends essentially all of its attention on LCD1 (aka 'classic' lattice). The LCD pathologic process involves the deposition of amyloid in the cornea, mainly in the mid- and anterior stroma.





What's the diagnosis?

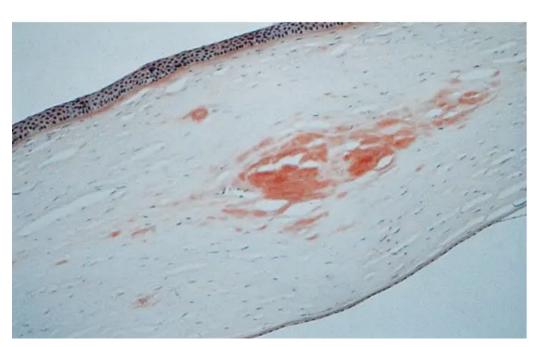
What's going on here? Again, let's first note what looks OK:

- --The epithelium seems to be unaffected --It's hard to see Bowman's the whole way across, and small sections may be hinky. But for the most part it appears intact—certainly lacking evidence of the extensive disruption we'd expect from a CDB or keratoconus. Now the wonky:
- --Stromal deposits, mainly in the central and anterior regions, are taking a stain.
- --The expected stromal clefts are missing around a glom of material, indicating distortion of the adjacent cornea's lamellar structure.

 This may be presented concurrently—the same slice of cornea under a different illumination in which the material glows green. The presence of anterior/mid-stromal material that stains red but glows green can be only one thing:

Lattice corneal dystrophy (LCD) is one of the *epithelial-stromal TGFBI* corneal dystrophies. The *BCSC* recognizes five variants, but spends essentially all of its attention on LCD1 (aka 'classic' lattice). The LCD pathologic process involves the deposition of amyloid in the cornea, mainly in the mid- and anterior stroma. The subepithelial/Bowman's portion be involved.





What's the diagnosis?

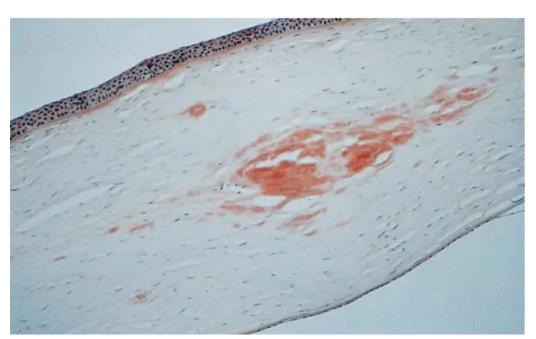
What's going on here? Again, let's first note what looks OK:

- --The epithelium seems to be unaffected --It's hard to see Bowman's the whole way across, and small sections may be hinky. But for the most part it appears intact—certainly lacking evidence of the extensive disruption we'd expect from a CDB or keratoconus. Now the wonky:
- --Stromal deposits, mainly in the central and anterior regions, are taking a stain.
- --The expected stromal clefts are missing around a glom of material, indicating distortion of the adjacent cornea's lamellar structure.

 This may be presented concurrently—the same slice of cornea under a different illumination in which the material glows green. The presence of anterior/mid-stromal material that stains red but glows green can be only one thing:

Lattice corneal dystrophy (LCD) is one of the *epithelial-stromal TGFBI* corneal dystrophies. The *BCSC* recognizes five variants, but spends essentially all of its attention on LCD1 (aka 'classic' lattice). The LCD pathologic process involves the deposition of amyloid in the cornea, mainly in the mid- and anterior stroma. The subepithelial/Bowman's portion may be involved.





What's the diagnosis?

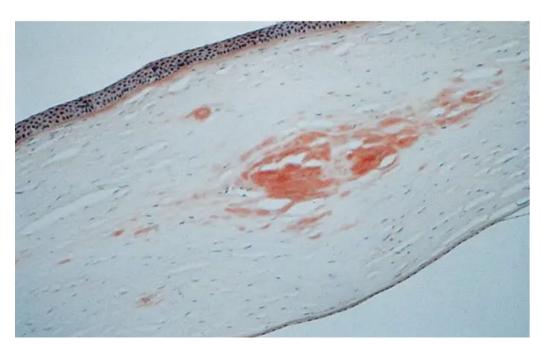
What's going on here? Again, let's first note what looks OK:

- --The epithelium seems to be unaffected
 --It's hard to see Bowman's the whole way
 across, and small sections may be hinky. But
 for the most part it appears intact—certainly
 lacking evidence of the extensive disruption
 we'd expect from a CDB or keratoconus.
 Now the wonky:
- --Stromal deposits, mainly in the central and anterior regions, are taking a stain.
- --The expected stromal clefts are missing around a glom of material, indicating distortion of the adjacent cornea's lamellar structure.

 This may be presented concurrently—the same slice of cornea under a different illumination in which the material glows green. The presence of anterior/mid-stromal material that stains red but glows green can be only one thing:

Lattice corneal dystrophy (LCD) is one of the *epithelial-stromal TGFBI* corneal dystrophies. The *BCSC* recognizes five variants, but spends essentially all of its attention on LCD1 (aka 'classic' lattice). The LCD pathologic process involves the deposition of amyloid in the cornea, mainly in the mid- and anterior stroma. The subepithelial/Bowman's portion may be involved. The classic stain (there are others) is





What's the diagnosis?

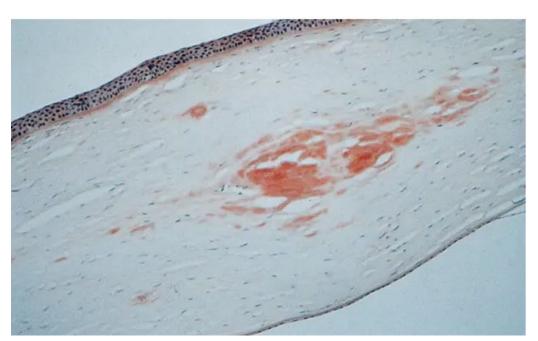
What's going on here? Again, let's first note what looks OK:

- --The epithelium seems to be unaffected
 --It's hard to see Bowman's the whole way
 across, and small sections may be hinky. But
 for the most part it appears intact—certainly
 lacking evidence of the extensive disruption
 we'd expect from a CDB or keratoconus.
 Now the wonky:
- --Stromal deposits, mainly in the central and anterior regions, are taking a stain.
- --The expected stromal clefts are missing around a glom of material, indicating distortion of the adjacent cornea's lamellar structure.

 This may be presented concurrently—the same slice of cornea under a different illumination in which the material glows green. The presence of anterior/mid-stromal material that stains red but glows green can be only one thing:

Lattice corneal dystrophy (LCD) is one of the *epithelial-stromal TGFBI* corneal dystrophies. The *BCSC* recognizes five variants, but spends essentially all of its attention on LCD1 (aka 'classic' lattice). The LCD pathologic process involves the deposition of amyloid in the cornea, mainly in the mid- and anterior stroma. The subepithelial/Bowman's portion may be involved. The classic stain (there are others) is Congo red





What's the diagnosis?

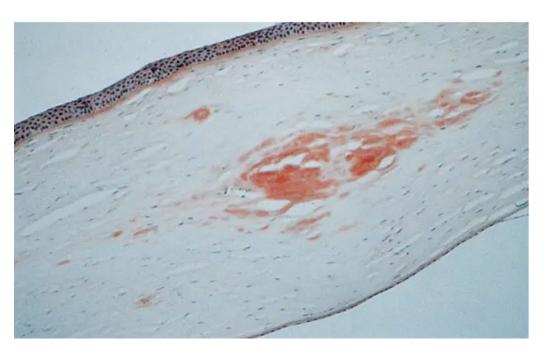
What's going on here? Again, let's first note what looks OK:

- --The epithelium seems to be unaffected
 --It's hard to see Bowman's the whole way
 across, and small sections may be hinky. But
 for the most part it appears intact—certainly
 lacking evidence of the extensive disruption
 we'd expect from a CDB or keratoconus.
 Now the wonky:
- --Stromal deposits, mainly in the central and anterior regions, are taking a stain.
- --The expected stromal clefts are missing around a glom of material, indicating distortion of the adjacent cornea's lamellar structure.

 This may be presented concurrently—the same slice of cornea under a different illumination in which the material glows green. The presence of anterior/mid-stromal material that stains red but glows green can be only one thing:

Lattice corneal dystrophy (LCD) is one of the *epithelial-stromal TGFBI* corneal dystrophies. The *BCSC* recognizes five variants, but spends essentially all of its attention on LCD1 (aka 'classic' lattice). The LCD pathologic process involves the deposition of amyloid in the cornea, mainly in the mid- and anterior stroma. The subepithelial/Bowman's portion may be involved. The classic stain (there are others) is Congo red; this is the stain that glows adj. -green when viewed with light.





What's the diagnosis?

What's going on here? Again, let's first note what looks OK:

- --The epithelium seems to be unaffected --It's hard to see Bowman's the whole way across, and small sections may be hinky. But for the most part it appears intact—certainly lacking evidence of the extensive disruption we'd expect from a CDB or keratoconus. Now the wonky:
- --Stromal deposits, mainly in the central and anterior regions, are taking a stain.
- --The expected stromal clefts are missing around a glom of material, indicating distortion of the adjacent cornea's lamellar structure.

 This may be presented concurrently—the same slice of cornea under a different illumination in which the material glows green. The presence of anterior/mid-stromal material that stains red but glows green can be only one thing:

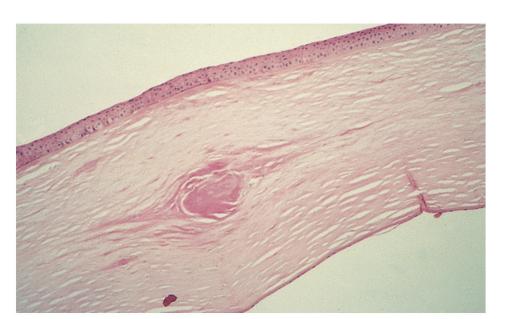
Lattice corneal dystrophy (LCD) is one of the *epithelial-stromal TGFBI* corneal dystrophies. The *BCSC* recognizes five variants, but spends essentially all of its attention on LCD1 (aka 'classic' lattice). The LCD pathologic process involves the deposition of amyloid in the cornea, mainly in the mid- and anterior stroma. The subepithelial/Bowman's portion may be involved. The classic stain (there are others) is Congo red; this is the stain that glows apple-green when viewed with polarized light.

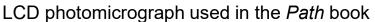


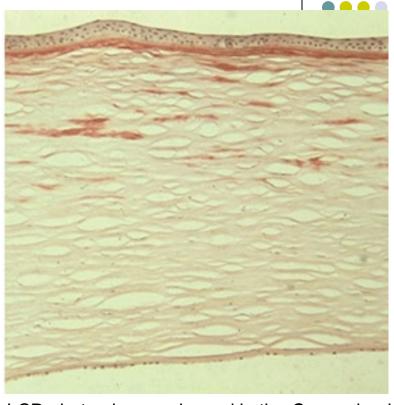


LCD photomicrograph used in the *Path* book

Note: The *Path* book asserts that the amyloid deposits are "fusiform" in distribution, a claim supported by the photomicrograph used in that volume (above).





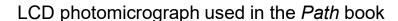


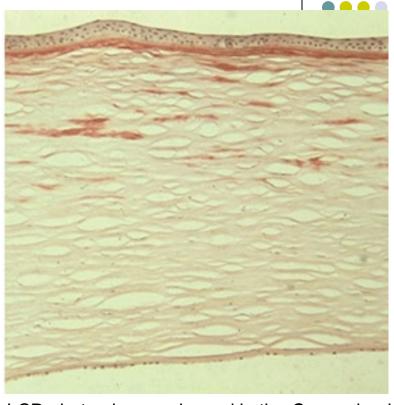
492

LCD photomicrograph used in the *Cornea* book

Note: The *Path* book asserts that the amyloid deposits are "fusiform" in distribution, a claim supported by the photomicrograph used in that volume (above). In contrast, the *Cornea* book does not use 'fusiform' in its description of LCD, and the amyloid deposits in the photomicrograph it uses do not demonstrate this quality (above, *right*).





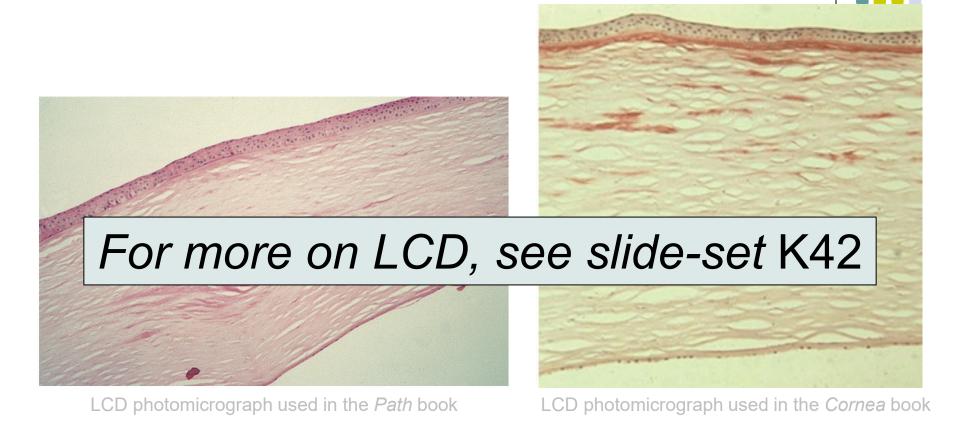


493

LCD photomicrograph used in the *Cornea* book

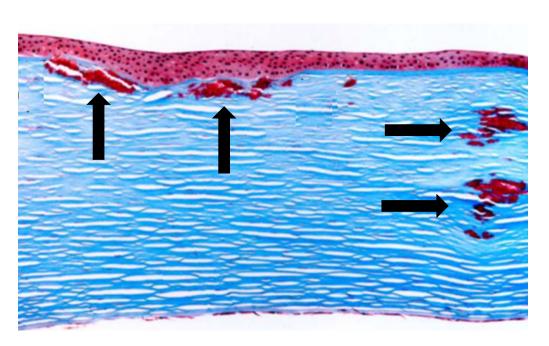
Note: The *Path* book asserts that the amyloid deposits are "fusiform" in distribution, a claim supported by the photomicrograph used in that volume (above). In contrast, the *Cornea* book does not use 'fusiform' in its description of LCD, and the amyloid deposits in the photomicrograph it uses do not demonstrate this quality (above, *right*). Which description is the one you should remember? I dunno. Caveat emptor.

494



Note: The *Path* book asserts that the amyloid deposits are "fusiform" in distribution, a claim supported by the photomicrograph used in that volume (above). In contrast, the *Cornea* book does not use 'fusiform' in its description of LCD, and the amyloid deposits in the photomicrograph it uses do not demonstrate this quality (above, *right*). Which description is the one you should remember? I dunno. Caveat emptor.

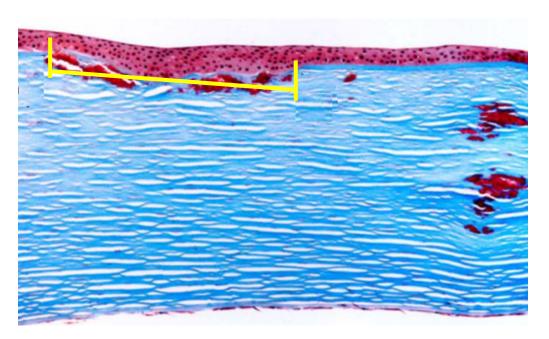




What's going on here?

--Stain-avid stromal deposits are present in the mid- and anterior stroma. Nearby clefts are compressed/distorted.

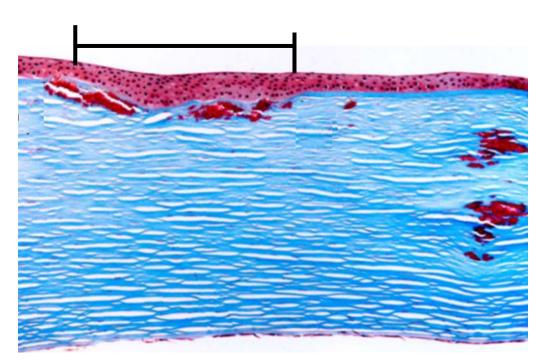




What's going on here?

- --Stain-avid stromal deposits are present in the mid- and anterior stroma. Nearby clefts are compressed/distorted.
- --A section of Bowman's is disrupted in association with subjacent deposits.

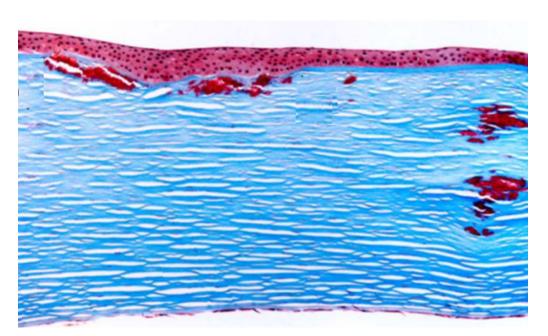




What's going on here?

- --Stain-avid stromal deposits are present in the mid- and anterior stroma. Nearby clefts are compressed/distorted.
- --A section of Bowman's is disrupted in association with subjacent deposits.
- --The epithelium is largely unaffected, with the notable exception of the portion associated with Bowman's disruption.





What's going on here?

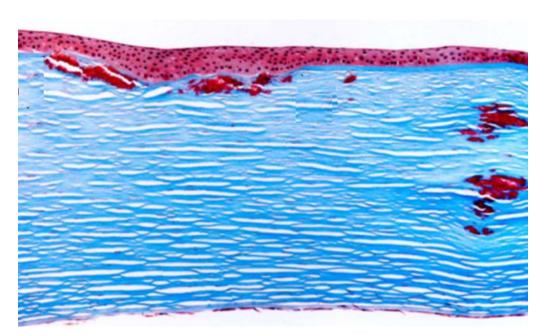
- --Stain-avid stromal deposits are present in the mid- and anterior stroma. Nearby clefts are compressed/distorted.
- --A section of Bowman's is disrupted in association with subjacent deposits.
- --The epithelium is largely unaffected, with the notable exception of the portion associated with Bowman's disruption.

This would be a tough one to get unless you're told the stain—

two words

.



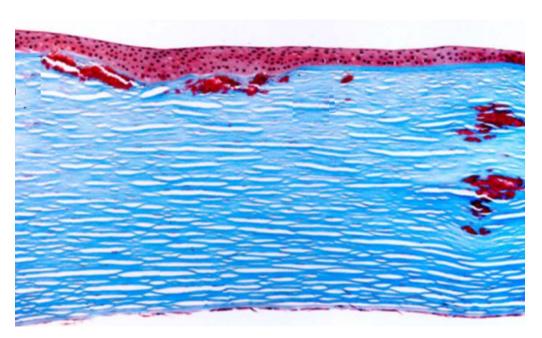


What's going on here?

- --Stain-avid stromal deposits are present in the mid- and anterior stroma. Nearby clefts are compressed/distorted.
- --A section of Bowman's is disrupted in association with subjacent deposits.
- --The epithelium is largely unaffected, with the notable exception of the portion associated with Bowman's disruption.

This would be a tough one to get unless you're told the stain—Masson trichrome.





What's the diagnosis?

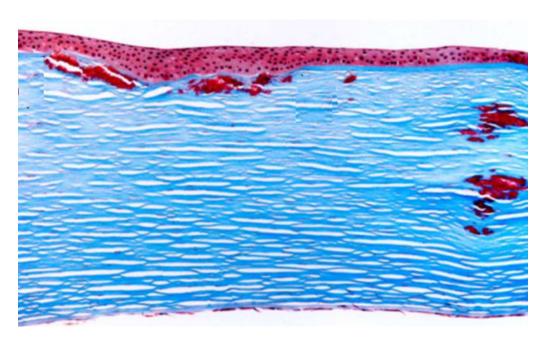
What's going on here?

- --Stain-avid stromal deposits are present in the mid- and anterior stroma. Nearby clefts are compressed/distorted.
- --A section of Bowman's is disrupted in association with subjacent deposits.
- --The epithelium is largely unaffected, with the notable exception of the portion associated with Bowman's disruption.

This would be a tough one to get unless you're told the stain—Masson trichrome.

The presence of anterior/mid-stromal deposits that stains avidly with Masson trichrome can be only one thing:





What's going on here?

- --Stain-avid stromal deposits are present in the mid- and anterior stroma. Nearby clefts are compressed/distorted.
- --A section of Bowman's is disrupted in association with subjacent deposits.
- --The epithelium is largely unaffected, with the notable exception of the portion associated with Bowman's disruption.

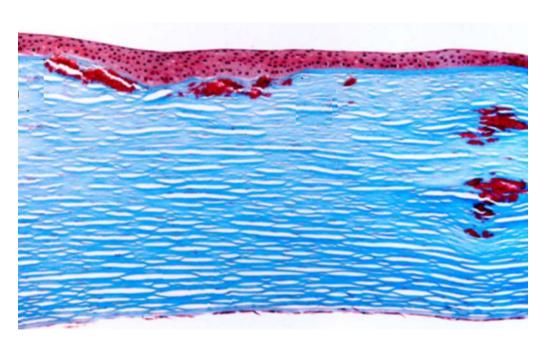
This would be a tough one to get unless you're told the stain—Masson trichrome.

The presence of anterior/mid-stromal deposits that stains avidly with Masson trichrome can be only one thing:

What's the diagnosis?

Granular corneal dystrophy type 1 (GCD1) one of the epithelial-stromal TGFBI corneal dystrophies.





What's going on here?

- --Stain-avid stromal deposits are present in the mid- and anterior stroma. Nearby clefts are compressed/distorted.
- --A section of Bowman's is disrupted in association with subjacent deposits.
- --The epithelium is largely unaffected, with the notable exception of the portion associated with Bowman's disruption.

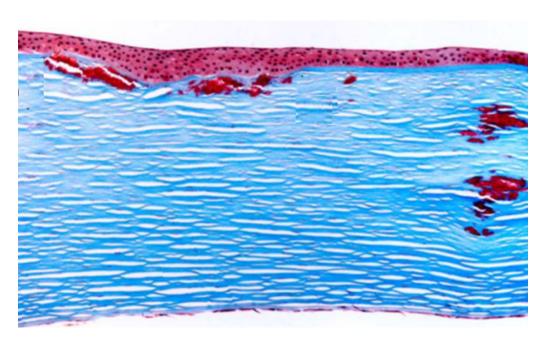
This would be a tough one to get unless you're told the stain—Masson trichrome.

The presence of anterior/mid-stromal deposits that stains avidly with Masson trichrome can be only one thing:

What's the diagnosis?

Granular corneal dystrophy type 1 (GCD1) is one of the epithelial-stromal TGFBI corneal dystrophies.





What's going on here?

- --Stain-avid stromal deposits are present in the mid- and anterior stroma. Nearby clefts are compressed/distorted.
- --A section of Bowman's is disrupted in association with subjacent deposits.
- --The epithelium is largely unaffected, with the notable exception of the portion associated with Bowman's disruption.

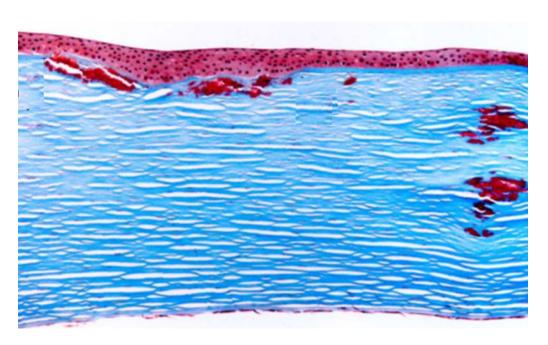
This would be a tough one to get unless you're told the stain—Masson trichrome.

The presence of anterior/mid-stromal deposits that stains avidly with Masson trichrome can be only one thing:

What's the diagnosis?

Granular corneal dystrophy type 1 (GCD1) is one of the *epithelial-stromal TGFBI* corneal dystrophies. The GCD1 pathologic process involves the deposition of in the mid- and anterior stroma.





What's going on here?

- --Stain-avid stromal deposits are present in the mid- and anterior stroma. Nearby clefts are compressed/distorted.
- --A section of Bowman's is disrupted in association with subjacent deposits.
- --The epithelium is largely unaffected, with the notable exception of the portion associated with Bowman's disruption.

This would be a tough one to get unless you're told the stain—Masson trichrome.

The presence of anterior/mid-stromal deposits that stains avidly with Masson trichrome can be only one thing:

What's the diagnosis?

Granular corneal dystrophy type 1 (GCD1) is one of the *epithelial-stromal TGFBI* corneal dystrophies. The GCD1 pathologic process involves the deposition of hyaline in the mid- and anterior stroma.



505

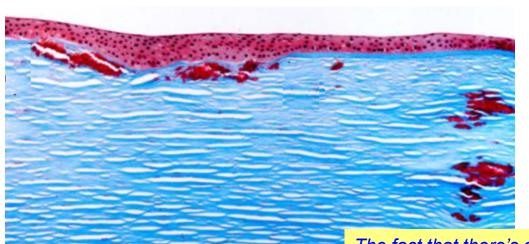
FYI: This is the GCD1 Masson trichrome photomicrograph used in the *Path* book.



FYI: This is the GCD1 Masson trichrome photomicrograph used in the *Path* book. Its caption: "Masson trichrome. The stromal collagen stains **blue**, and the granular hyaline deposits stains **brilliant red**."







What's going on here?

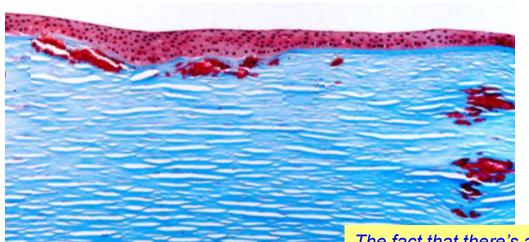
- --Stain-avid stromal deposits are present in the mid- and anterior stroma. Nearby clefts are compressed/distorted.
- --A section of Bowman's is disrupted in association with subjacent deposits.
- --The epithelium is largely unaffected, with the notable exception of the portion associated with Bowman's disruption.

This would be a tough one to get unless you're

The fact that there's a 'type 1' GCD implies the existence of a type 2 (at the very least). Is this the case?

Granular corneal dystrophy type 1





What's going on here?

- --Stain-avid stromal deposits are present in the mid- and anterior stroma. Nearby clefts are compressed/distorted.
- --A section of Bowman's is disrupted in association with subjacent deposits.
- --The epithelium is largely unaffected, with the notable exception of the portion associated with Bowman's disruption.

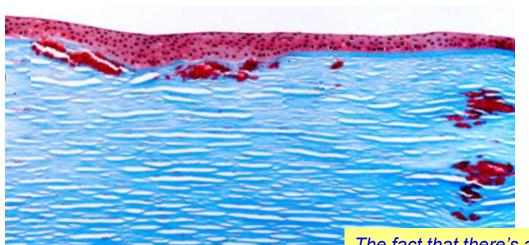
This would be a tough one to get unless you're

The fact that there's a 'type 1' GCD implies the existence of a type 2 (at the very least). Is this the case?

Indeed it is—granular corneal dystrophy type 2 (GCD2) is a thing

Granular corneal dystrophy type 1





What's going on here?

- --Stain-avid stromal deposits are present in the mid- and anterior stroma. Nearby clefts are compressed/distorted.
- --A section of Bowman's is disrupted in association with subjacent deposits.
- --The epithelium is largely unaffected, with the notable exception of the portion associated with Bowman's disruption.

This would be a tough one to get unless you're

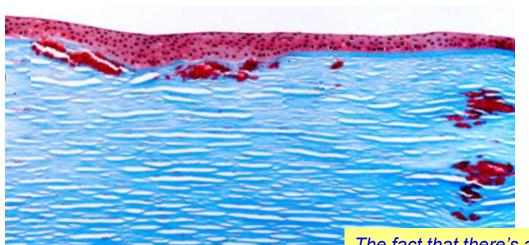
The fact that there's a 'type 1' GCD implies the existence of a type 2 (at the very least). Is this the case?

Indeed it is—granular corneal dystrophy type 2 (GCD2) is a thing

GCD2 also has an eponymous name—what is it?

Granular corneal dystrophy type 1





What's going on here?

- --Stain-avid stromal deposits are present in the mid- and anterior stroma. Nearby clefts are compressed/distorted.
- --A section of Bowman's is disrupted in association with subjacent deposits.
- --The epithelium is largely unaffected, with the notable exception of the portion associated with Bowman's disruption.

This would be a tough one to get unless you're

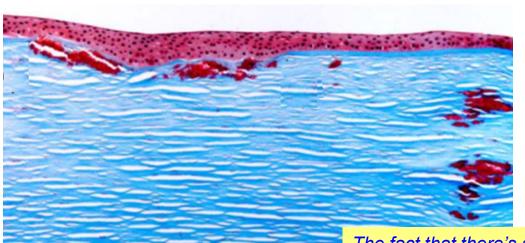
The fact that there's a 'type 1' GCD implies the existence of a type 2 (at the very least). Is this the case?

Indeed it is—granular corneal dystrophy type 2 (GCD2) is a thing

GCD2 also has an eponymous name—what is it? Avellino corneal dystrophy

Granular corneal dystrophy type 1





What's going on here?

- --Stain-avid stromal deposits are present in the mid- and anterior stroma. Nearby clefts are compressed/distorted.
- --A section of Bowman's is disrupted in association with subjacent deposits.
- --The epithelium is largely unaffected, with the notable exception of the portion associated with Bowman's disruption.

This would be a tough one to get unless you're

The fact that there's a 'type 1' GCD implies the existence of a type 2 (at the very least). Is this the case?

Indeed it is—granular corneal dystrophy type 2 (GCD2) is a thing

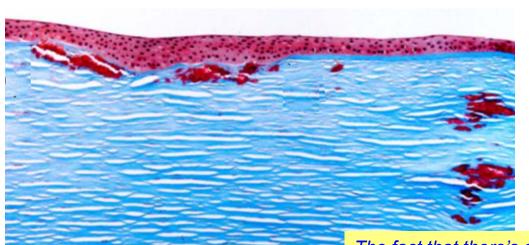
GCD2 also has an eponymous name—what is it? Avellino corneal dystrophy

Granular corneal dystrophy type 1

The GCD1 pathologic process involves

In a nutshell, what is GCD2?





What's going on here?

- --Stain-avid stromal deposits are present in the mid- and anterior stroma. Nearby clefts are compressed/distorted.
- -- A section of Bowman's is disrupted in association with subjacent deposits.
- -- The epithelium is largely unaffected, with the notable exception of the portion associated with Bowman's disruption.

This would be a tough one to get unless you're

The fact that there's a 'type 1' GCD implies the existence of a type 2 (at the very least). Is this the case?

Indeed it is—granular corneal dystrophy type 2 (GCD2) is a thing

GCD2 also has an eponymous name—what is it? Avellino corneal dystrophy

Granular corneal dystrophy type 1

GCD1 pathologic process in

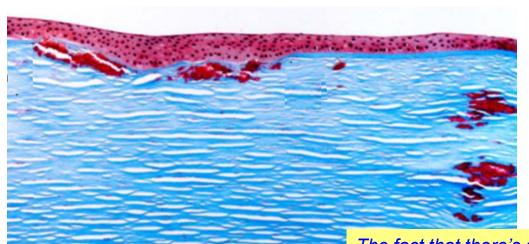
In a nutshell, what is GCD2?

abb.

Both clinically and histopathologically, it is a combo of abb.

land





What's going on here?

- --Stain-avid stromal deposits are present in the mid- and anterior stroma. Nearby clefts are compressed/distorted.
- --A section of Bowman's is disrupted in association with subjacent deposits.
- --The epithelium is largely unaffected, with the notable exception of the portion associated with Bowman's disruption.

This would be a tough one to get unless you're

The fact that there's a 'type 1' GCD implies the existence of a type 2 (at the very least). Is this the case?

Indeed it is—granular corneal dystrophy type 2 (GCD2) is a thing

GCD2 also has an eponymous name—what is it? Avellino corneal dystrophy

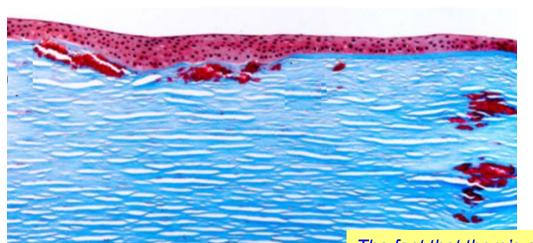
Granular corneal dystrophy type 1

The GCD1 pathologic process involves

In a nutshell, what is GCD2?

Both clinically and histopathologically, it is a combo of LCD and GCD1.





What's going on here?

- --Stain-avid stromal deposits are present in the mid- and anterior stroma. Nearby clefts are compressed/distorted.
- --A section of Bowman's is disrupted in association with subjacent deposits.
- --The epithelium is largely unaffected, with the notable exception of the portion associated with Bowman's disruption.

This would be a tough one to get unless you're

The fact that there's a 'type 1' GCD implies the existence of a type 2 (at the very least). Is this the case?

Indeed it is—granular corneal dystrophy type 2 (GCD2) is a thing

GCD2 also has an eponymous name—what is it? Avellino corneal dystrophy

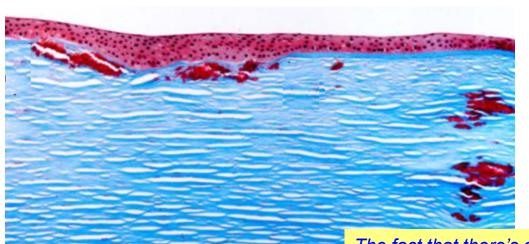
Granular corneal dystrophy type 1

The GCD1 pathologic process involves

In a nutshell, what is GCD2?

Both clinically and histopathologically, it is a combo of LCD and GCD1. As would be expected, a GCD2 cornea contains deposits of both and





What's going on here?

- --Stain-avid stromal deposits are present in the mid- and anterior stroma. Nearby clefts are compressed/distorted.
- --A section of Bowman's is disrupted in association with subjacent deposits.
- --The epithelium is largely unaffected, with the notable exception of the portion associated with Bowman's disruption.

This would be a tough one to get unless you're

The fact that there's a 'type 1' GCD implies the existence of a type 2 (at the very least). Is this the case?

Indeed it is—granular corneal dystrophy type 2 (GCD2) is a thing

GCD2 also has an eponymous name—what is it? Avellino corneal dystrophy

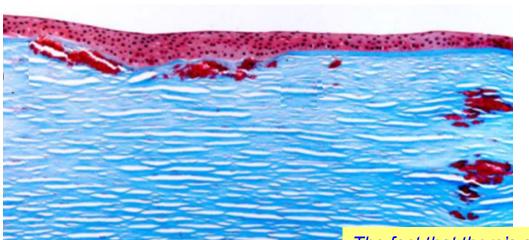
Granular corneal dystrophy type 1

The GCD1 pathologic process involves

In a nutshell, what is GCD2?

Both clinically and histopathologically, it is a combo of LCD and GCD1. As would be expected, a GCD2 cornea contains deposits of both amyloid and hyaline





What's going on here?

- --Stain-avid stromal deposits are present in the mid- and anterior stroma. Nearby clefts are compressed/distorted.
- --A section of Bowman's is disrupted in association with subjacent deposits.
- --The epithelium is largely unaffected, with the notable exception of the portion associated with Bowman's disruption.

This would be a tough one to get unless you're

The fact that there's a 'type 1' GCD implies the existence of a type 2 (at the very least). Is this the case?

Indeed it is—granular corneal dystrophy type 2 (GCD2) is a thing

GCD2 also has an eponymous name—what is it? Avellino corneal dystrophy

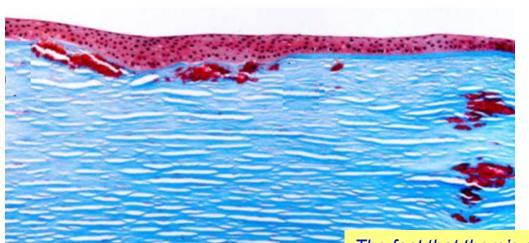
Granular corneal dystrophy type 1

The GCD1 pathologic process involves

In a nutshell, what is GCD2?

Both clinically and histopathologically, it is a combo of LCD and GCD1. As would be expected, a GCD2 cornea contains deposits of both amyloid and hyaline, and thus stains with two words and both.





What's going on here?

- --Stain-avid stromal deposits are present in the mid- and anterior stroma. Nearby clefts are compressed/distorted.
- --A section of Bowman's is disrupted in association with subjacent deposits.
- --The epithelium is largely unaffected, with the notable exception of the portion associated with Bowman's disruption.

This would be a tough one to get unless you're

The fact that there's a 'type 1' GCD implies the existence of a type 2 (at the very least). Is this the case?

Indeed it is—granular corneal dystrophy type 2 (GCD2) is a thing

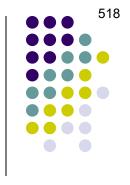
GCD2 also has an eponymous name—what is it? Avellino corneal dystrophy

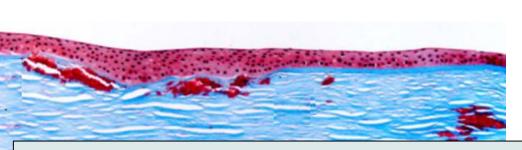
Granular corneal dystrophy type 1

The GCD1 pathologic process involves

In a nutshell, what is GCD2?

Both clinically and histopathologically, it is a combo of LCD and GCD1. As would be expected, a GCD2 cornea contains deposits of both amyloid and hyaline, and thus stains with Congo red and Masson trichrome both.





What's going on here?

- --Stain-avid stromal deposits are present in the mid- and anterior stroma. Nearby clefts are compressed/distorted.
- --A section of Bowman's is disrupted in association with subjacent deposits.

The continue in lease because if a steel with the

For more on GCD1&2, see slide-set K42

This would be a tough one to get unless you're

The fact that there's a 'type 1' GCD implies the existence of a type 2' (at the very least). Is this the case?

Indeed it is—granular corneal dystrophy type 2 (GCD2) is a thing

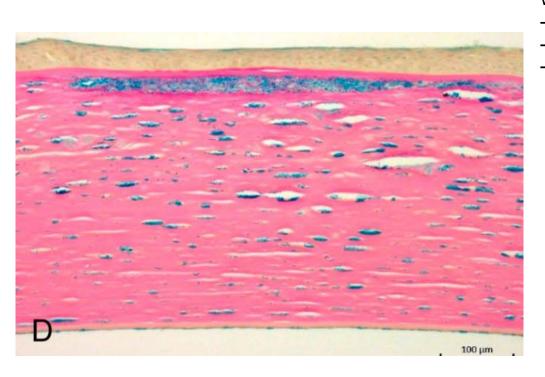
GCD2 also has an eponymous name—what is it? Avellino corneal dystrophy

Granular corneal dystrophy type 1 ()
The GCD1 pathologic process involves

In a nutshell, what is GCD2?

Both clinically and histopathologically, it is a combo of LCD and GCD1. As would be expected, a GCD2 cornea contains deposits of both amyloid and hyaline, and thus stains with Congo red and Masson trichrome both.





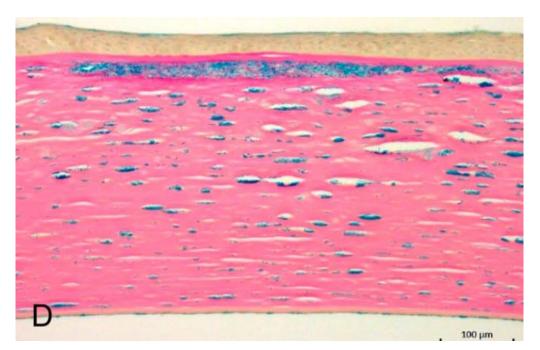
What's going on here?
--The epi looks involved vs not

--Bowman's seems --Stromal clefting is

affected vs not

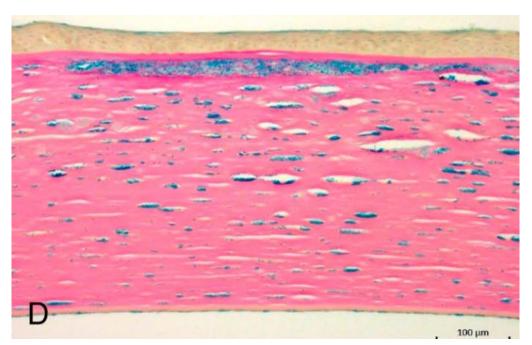
disrupted vs not





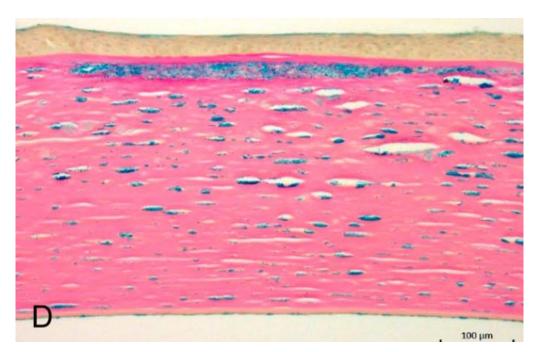
- --The epi looks OK
- --Bowman's seems intact
- --Stromal clefting is unremarkable





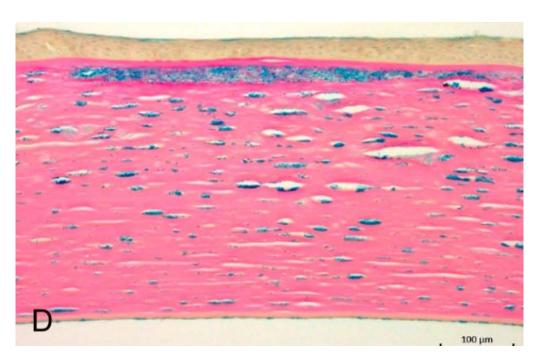
- --The epi looks OK
- --Bowman's seems intact
- --Stromal clefting is unremarkable But also:
- --Stain-avid deposits are present through the extent of the stroma.





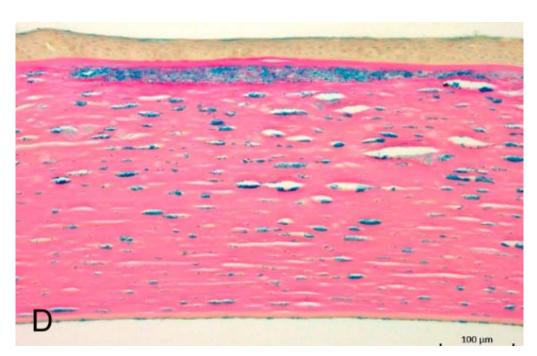
- --The epi looks OK
- --Bowman's seems intact
- --Stromal clefting is unremarkable But also:
- --Stain-avid deposits are present through the entire breadth of the stroma.





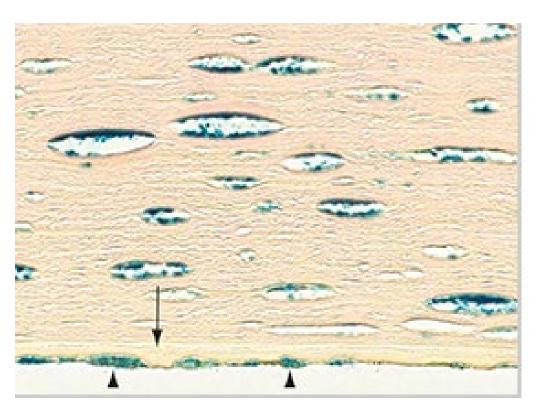
- --The epi looks OK
- --Bowman's seems intact
- --Stromal clefting is unremarkable But also:
- --Stain-avid deposits are present through the entire breadth of the stroma.
- --Descemet's and the endothelium are v involved.





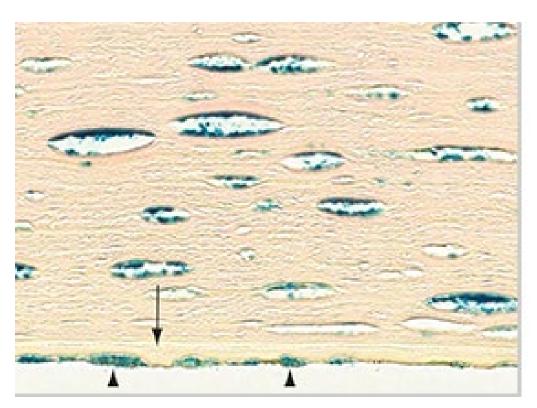
- --The epi looks OK
- --Bowman's seems intact
- --Stromal clefting is unremarkable But also:
- --Stain-avid deposits are present through the entire breadth of the stroma.
- --Descemet's and the endothelium are involved.





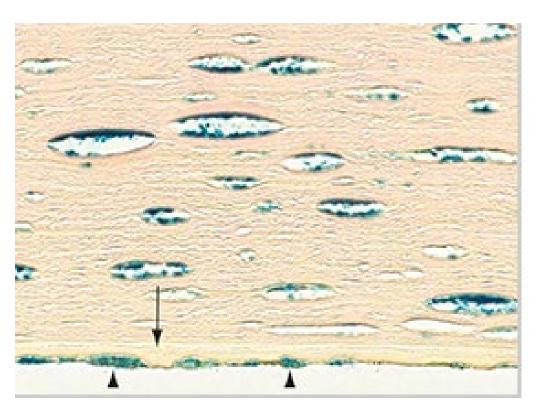
- --The epi looks OK
- --Bowman's seems intact
- --Stromal clefting is unremarkable But also:
- --Stain-avid deposits are present through the entire breadth of the stroma.
- --Descemet's and the endothelium are involved. This can be seen more easily **here**, where the *arrowheads* indicate stain in the





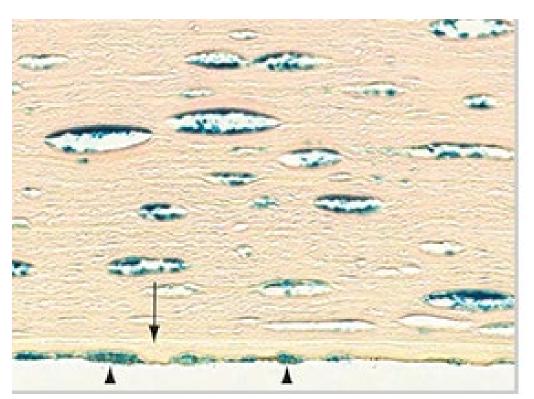
- --The epi looks OK
- --Bowman's seems intact
- --Stromal clefting is unremarkable But also:
- --Stain-avid deposits are present through the entire breadth of the stroma.
- --Descemet's and the endothelium are involved. This can be seen more easily **here**, where the *arrowheads* indicate stain in the endothelium





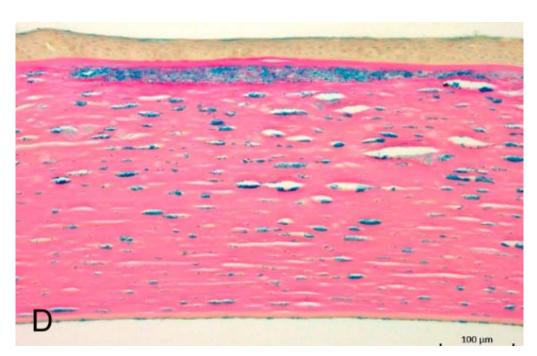
- --The epi looks OK
- --Bowman's seems intact
- --Stromal clefting is unremarkable But also:
- --Stain-avid deposits are present through the entire breadth of the stroma.





- --The epi looks OK
- --Bowman's seems intact
- --Stromal clefting is unremarkable But also:
- --Stain-avid deposits are present through the entire breadth of the stroma.
- --Descemet's and the endothelium are involved. This can be seen more easily **here**, where the *arrowheads* indicate stain in the endothelium, and the *arrow* is pointing to an excrescence in Descemet's membrane, ie, a *guttae*. (Note: This specimen used a different stain than the first.)



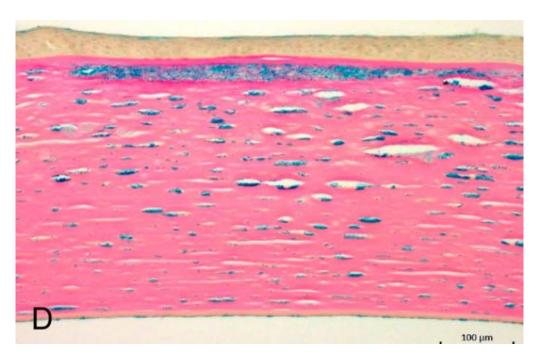


What's going on here?

- --The epi looks OK
- --Bowman's seems intact
- --Stromal clefting is unremarkable But also:
- --Stain-avid deposits are present through the entire breadth of the stroma.
- --Descemet's and the endothelium are involved. This can be seen more easily **here**, where the *arrowheads* indicate stain in the endothelium, and the *arrow* is pointing to an excrescence in Descemet's membrane, ie, a *guttae*. (Note: This specimen used a different stain than the first.)

The presence of deposits through the entire stroma and into Descemet's/endothelium can be only one thing:





What's going on here?

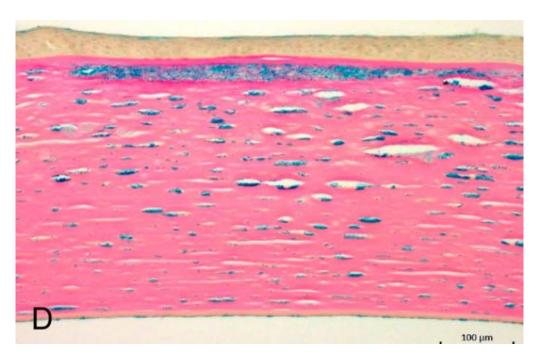
- --The epi looks OK
- --Bowman's seems intact
- --Stromal clefting is unremarkable But also:
- --Stain-avid deposits are present through the entire breadth of the stroma.
- --Descemet's and the endothelium are involved. This can be seen more easily **here**, where the *arrowheads* indicate stain in the endothelium, and the *arrow* is pointing to an excrescence in Descemet's membrane, ie, a *guttae*. (Note: This specimen used a different stain than the first.)

The presence of deposits through the entire stroma and into Descemet's/endothelium can be only one thing:

What's the diagnosis?

Macular corneal dystrophy (MCD) is v one of the epithelial-stromal TGFBI corneal dystrophies.





What's going on here?

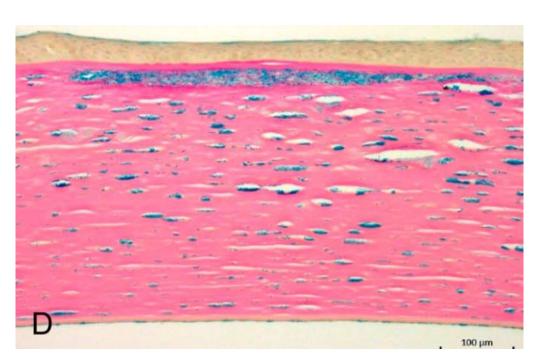
- --The epi looks OK
- --Bowman's seems intact
- --Stromal clefting is unremarkable But also:
- --Stain-avid deposits are present through the entire breadth of the stroma.
- --Descemet's and the endothelium are involved. This can be seen more easily **here**, where the *arrowheads* indicate stain in the endothelium, and the *arrow* is pointing to an excrescence in Descemet's membrane, ie, a *guttae*. (Note: This specimen used a different stain than the first.)

The presence of deposits through the entire stroma and into Descemet's/endothelium can be only one thing:

What's the diagnosis?

Macular corneal dystrophy (MCD) isn't one of the epithelial-stromal TGFBI corneal dystrophies.





What's going on here?

- --The epi looks OK
- --Bowman's seems intact
- --Stromal clefting is unremarkable But also:
- --Stain-avid deposits are present through the entire breadth of the stroma.
- --Descemet's and the endothelium are involved. This can be seen more easily **here**, where the *arrowheads* indicate stain in the endothelium, and the *arrow* is pointing to an excrescence in Descemet's membrane, ie, a *guttae*. (Note: This specimen used a different stain than the first.)

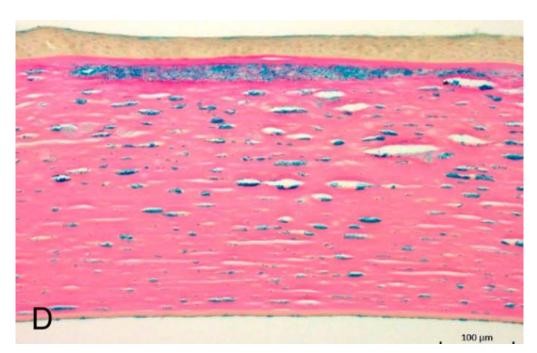
The presence of deposits through the entire stroma and into Descemet's/endothelium can be only one thing:

What's the diagnosis?

Macular corneal dystrophy (MCD) isn't one of the epithelial-stromal TGFBI corneal dystrophies.

The abnormal material deposited in the cornea is





What's going on here?

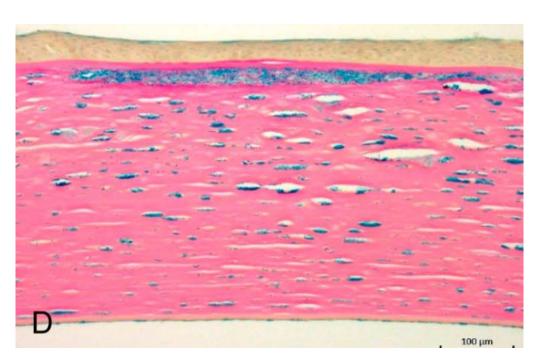
- --The epi looks OK
- --Bowman's seems intact
- --Stromal clefting is unremarkable But also:
- --Stain-avid deposits are present through the entire breadth of the stroma.
- --Descemet's and the endothelium are involved. This can be seen more easily **here**, where the *arrowheads* indicate stain in the endothelium, and the *arrow* is pointing to an excrescence in Descemet's membrane, ie, a *guttae*. (Note: This specimen used a different stain than the first.)

The presence of deposits through the entire stroma and into Descemet's/endothelium can be only one thing:

What's the diagnosis?

Macular corneal dystrophy (MCD) *isn't* one of the *epithelial-stromal TGFBI* corneal dystrophies. The abnormal material deposited in the cornea is glycosaminoglycans (GAGs).





What's going on here?

- --The epi looks OK
- --Bowman's seems intact
- --Stromal clefting is unremarkable But also:
- --Stain-avid deposits are present through the entire breadth of the stroma.
- --Descemet's and the endothelium are involved. This can be seen more easily **here**, where the *arrowheads* indicate stain in the endothelium, and the *arrow* is pointing to an excrescence in Descemet's membrane, ie, a *guttae*. (Note: This specimen used a different stain than the first.)

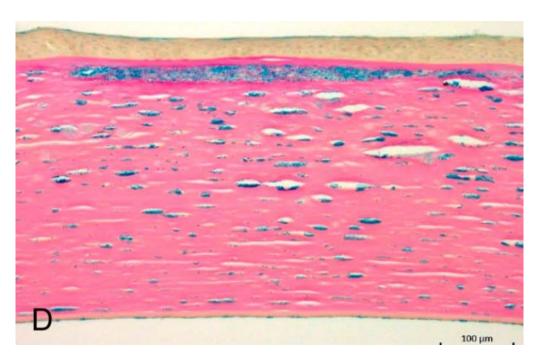
The presence of deposits through the entire stroma and into Descemet's/endothelium can be only one thing:

What's the diagnosis?

Macular corneal dystrophy (MCD) *isn't* one of the *epithelial-stromal TGFBI* corneal dystrophies. The abnormal material deposited in the cornea is glycosaminoglycans (GAGs).*

*Were you expecting here?





What's going on here?

- --The epi looks OK
- --Bowman's seems intact
- --Stromal clefting is unremarkable But also:
- --Stain-avid deposits are present through the entire breadth of the stroma.
- --Descemet's and the endothelium are involved. This can be seen more easily **here**, where the *arrowheads* indicate stain in the endothelium, and the *arrow* is pointing to an excrescence in Descemet's membrane, ie, a *guttae*. (Note: This specimen used a different stain than the first.)

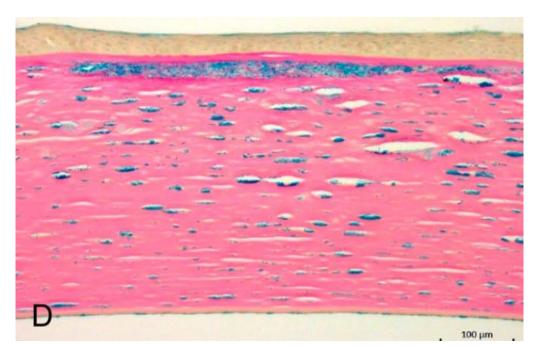
The presence of deposits through the entire stroma and into Descemet's/endothelium can be only one thing:

What's the diagnosis?

Macular corneal dystrophy (MCD) *isn't* one of the *epithelial-stromal TGFBI* corneal dystrophies. The abnormal material deposited in the cornea is glycosaminoglycans (GAGs) .*

*Were you expecting *mucopolysaccharides* here?





What's going on here?

- --The epi looks OK
- --Bowman's seems intact
- --Stromal clefting is unremarkable But also:
- --Stain-avid deposits are present through the entire breadth of the stroma.
- --Descemet's and the endothelium are involved. This can be seen more easily **here**, where the *arrowheads* indicate stain in the endothelium, and the *arrow* is pointing to an excrescence in Descemet's membrane, ie, a *guttae*. (Note: This specimen used a different stain than the first.)

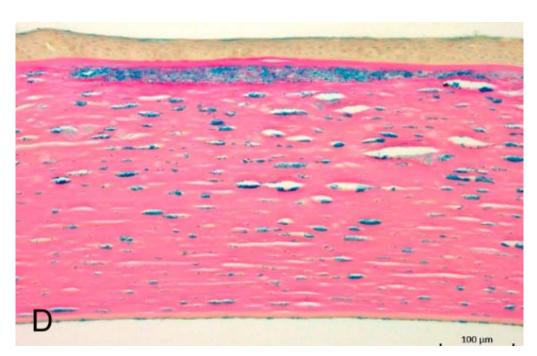
The presence of deposits through the entire stroma and into Descemet's/endothelium can be only one thing:

What's the diagnosis?

Macular corneal dystrophy (MCD) *isn't* one of the *epithelial-stromal TGFBI* corneal dystrophies. The abnormal material deposited in the cornea is glycosaminoglycans (GAGs).*

*Were you expecting *mucopolysaccharides* here? While not wrong, this term has been largely supplanted by *GAG* in the *BCSC*, so don't be confused if you see GAG on a test (and adjust your mnemonic accordingly, Marilyn).





What's going on here?

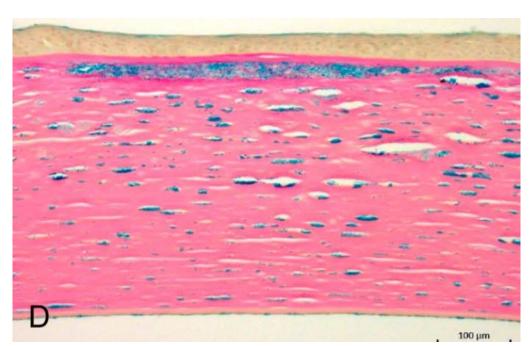
- --The epi looks OK
- --Bowman's seems intact
- --Stromal clefting is unremarkable But also:
- --Stain-avid deposits are present through the entire breadth of the stroma.
- --Descemet's and the endothelium are involved. This can be seen more easily **here**, where the *arrowheads* indicate stain in the endothelium, and the *arrow* is pointing to an excrescence in Descemet's membrane, ie, a *guttae*. (Note: This specimen used a different stain than the first.)

The presence of deposits through the entire stroma and into Descemet's/endothelium can be only one thing:

What's the diagnosis?

Macular corneal dystrophy (MCD) *isn't* one of the *epithelial-stromal TGFBI* corneal dystrophies. The abnormal material deposited in the cornea is glycosaminoglycans (GAGs).* The stain most closely associated with MCD is two words.





What's going on here?

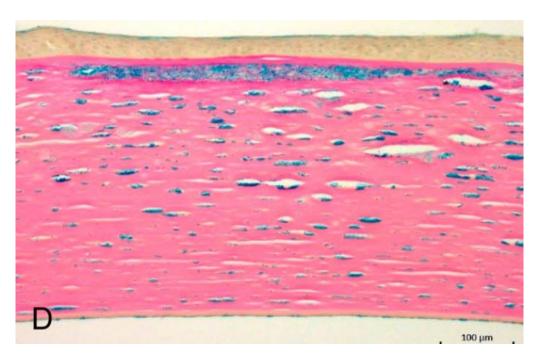
- --The epi looks OK
- --Bowman's seems intact
- --Stromal clefting is unremarkable But also:
- --Stain-avid deposits are present through the entire breadth of the stroma.
- --Descemet's and the endothelium are involved. This can be seen more easily **here**, where the *arrowheads* indicate stain in the endothelium, and the *arrow* is pointing to an excrescence in Descemet's membrane, ie, a *guttae*. (Note: This specimen used a different stain than the first.)

The presence of deposits through the entire stroma and into Descemet's/endothelium can be only one thing:

What's the diagnosis?

Macular corneal dystrophy (MCD) *isn't* one of the *epithelial-stromal TGFBI* corneal dystrophies. The abnormal material deposited in the cornea is glycosaminoglycans (GAGs).* The stain most closely associated with MCD is Alcian blue.





What's going on here?

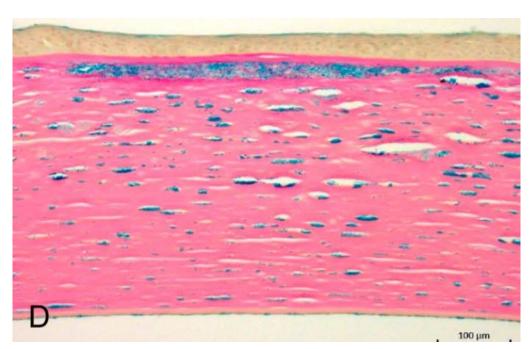
- --The epi looks OK
- --Bowman's seems intact
- --Stromal clefting is unremarkable But also:
- --Stain-avid deposits are present through the entire breadth of the stroma.
- --Descemet's and the endothelium are involved. This can be seen more easily **here**, where the *arrowheads* indicate stain in the endothelium, and the *arrow* is pointing to an excrescence in Descemet's membrane, ie, a *guttae*. (Note: This specimen used a different stain than the first.)

The presence of deposits through the entire stroma and into Descemet's/endothelium can be only one thing:

What's the diagnosis?

Macular corneal dystrophy (MCD) *isn't* one of the *epithelial-stromal TGFBI* corneal dystrophies. The abnormal material deposited in the cornea is glycosaminoglycans (GAGs).* The stain most closely associated with MCD is Alcian blue. Unlike vs As with MCD pts are not vote to the total dystrophies discussed previously, at risk for REE.





What's going on here?

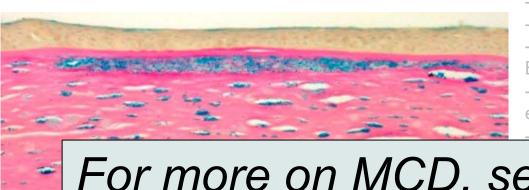
- --The epi looks OK
- --Bowman's seems intact
- --Stromal clefting is unremarkable But also:
- --Stain-avid deposits are present through the entire breadth of the stroma.
- --Descemet's and the endothelium are involved. This can be seen more easily **here**, where the *arrowheads* indicate stain in the endothelium, and the *arrow* is pointing to an excrescence in Descemet's membrane, ie, a *guttae*. (Note: This specimen used a different stain than the first.)

The presence of deposits through the entire stroma and into Descemet's/endothelium can be only one thing:

What's the diagnosis?

Macular corneal dystrophy (MCD) *isn't* one of the *epithelial-stromal TGFBI* corneal dystrophies. The abnormal material deposited in the cornea is glycosaminoglycans (GAGs).* The stain most closely associated with MCD is Alcian blue. Unlike the *TGFBI* dystrophies discussed previously, MCD pts are not at risk for REE.





What's going on here?

- --The epi looks OK
- --Bowman's seems intact
- --Stromal clefting is unremarkable But also:
- --Stain-avid deposits are present through the entire breadth of the stroma.

--Descemet's and the endothelium_are_involved.

For more on MCD, see slide-set K43

the ium , ce in



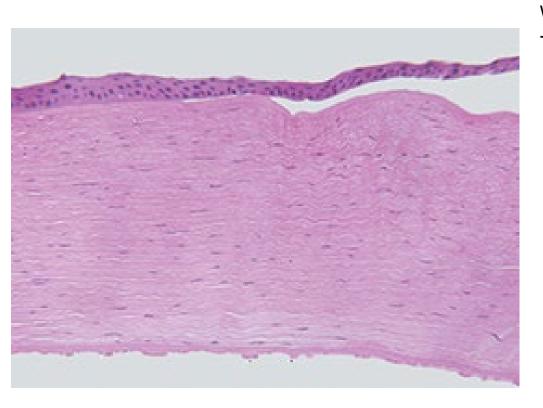
Descemet's membrane, ie, a *guttae*. (Note: This specimen used a different stain than the first.)

The presence of deposits through the entire stroma and into Descemet's/endothelium can be only one thing:

What's the diagnosis?

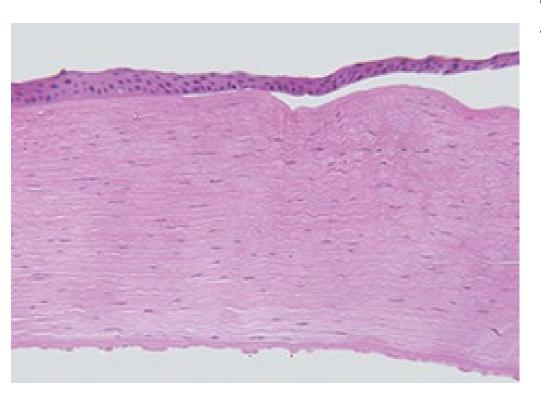
Macular corneal dystrophy (MCD) *isn't* one of the *epithelial-stromal TGFBI* corneal dystrophies. The abnormal material deposited in the cornea is glycosaminoglycans (GAGs).* The stain most closely associated with MCD is Alcian blue. Unlike the *TGFBI* dystrophies discussed previously, MCD pts are not at risk for REE.





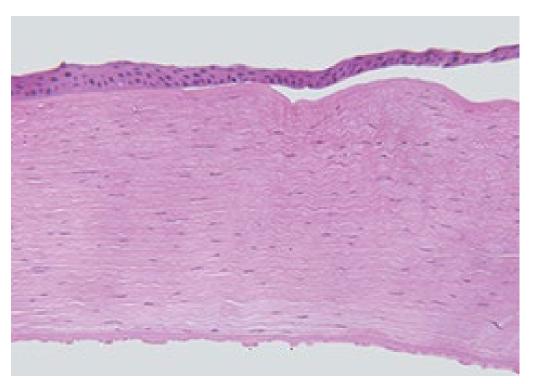
What's going on here?
--The epi looks involved vuninvolved vuninvolved





What's going on here? --The epi looks bad.

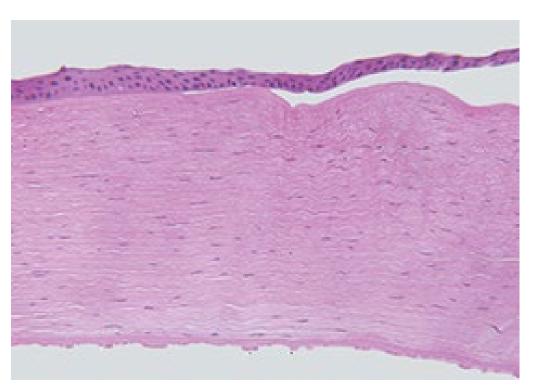




What's going on here?

--The epi looks bad—it's thin in places, and over half has separated from Bowman's.





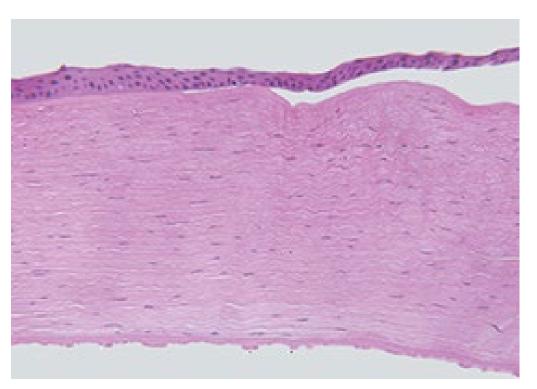
What's going on here?

--The epi looks bad—it's thin in places, and over half has separated from Bowman's.

--Speaking of: Bowman's seems

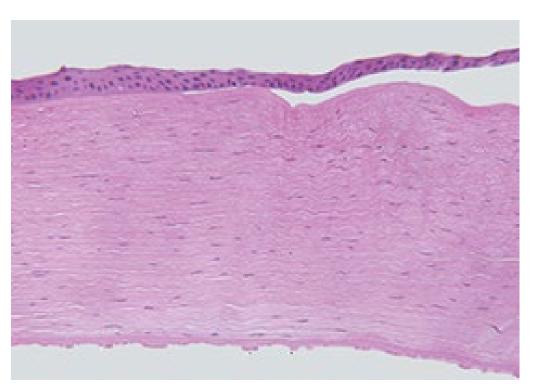






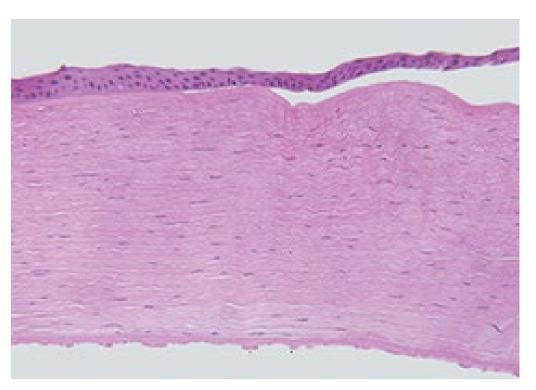
- --The epi looks bad—it's thin in places, and over half has separated from Bowman's.
- --Speaking of: Bowman's seems intact





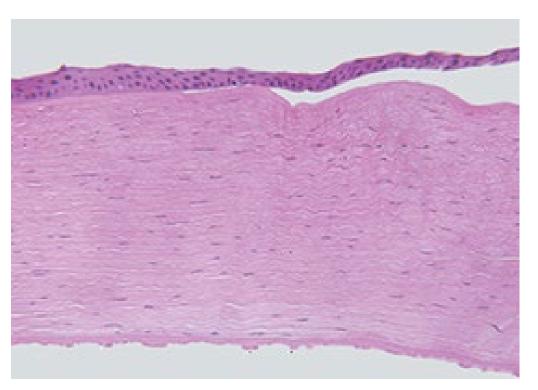
- --The epi looks bad—it's thin in places, and over half has separated from Bowman's.
- --Speaking of: Bowman's seems intact
- --The stroma is v taking a special stain, suggesting this is v a stromal dystrophy.





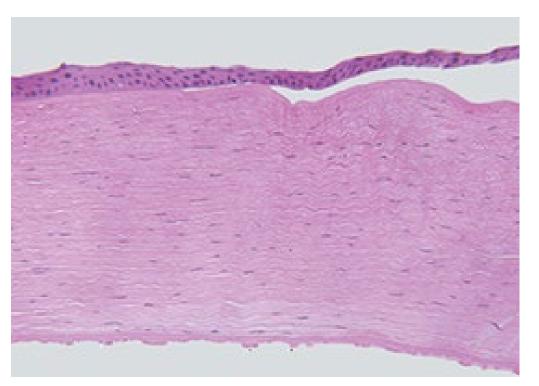
- --The epi looks bad—it's thin in places, and over half has separated from Bowman's.
- --Speaking of: Bowman's seems intact
- --The stroma isn't taking a special stain, suggesting this isn't a stromal dystrophy.





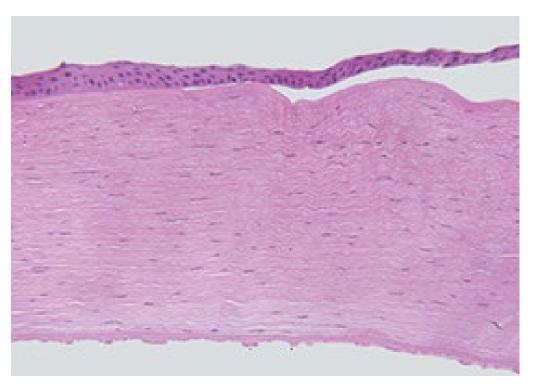
- --The epi looks bad—it's thin in places, and over half has separated from Bowman's.
- --Speaking of: Bowman's seems intact
- --The stroma isn't taking a special stain, suggesting this isn't a stromal dystrophy.
- --Speaking of: Stromal clefting is affected vs unaffected





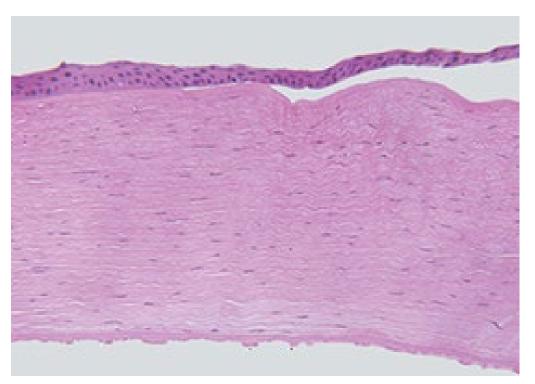
- --The epi looks bad—it's thin in places, and over half has separated from Bowman's.
- --Speaking of: Bowman's seems intact
- --The stroma isn't taking a special stain, suggesting this isn't a stromal dystrophy.
- --Speaking of: Stromal clefting is almost absent





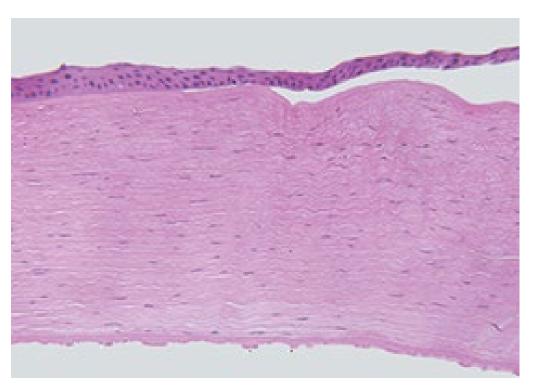
- --The epi looks bad—it's thin in places, and over half has separated from Bowman's.
- --Speaking of: Bowman's seems intact
- --The stroma isn't taking a special stain, suggesting this isn't a stromal dystrophy.
- --Speaking of: Stromal clefting is almost absent, suggesting and/or is present





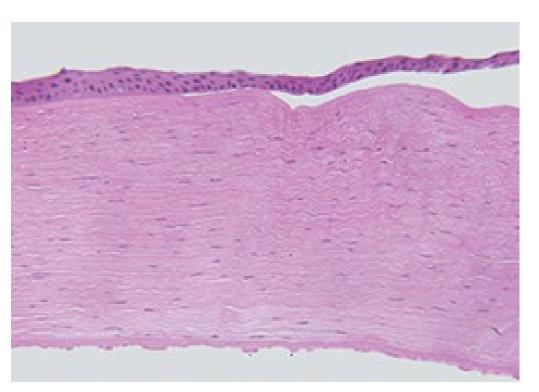
- --The epi looks bad—it's thin in places, and over half has separated from Bowman's.
- --Speaking of: Bowman's seems intact
- --The stroma isn't taking a special stain, suggesting this isn't a stromal dystrophy.
- --Speaking of: Stromal clefting is almost absent, suggesting scarring and/or edema is present





- --The epi looks bad—it's thin in places, and over half has separated from Bowman's.
- --Speaking of: Bowman's seems intact
- --The stroma isn't taking a special stain, suggesting this isn't a stromal dystrophy.
- --Speaking of: Stromal clefting is almost absent, suggesting scarring and/or edema is present Turning our attention to the posterior cornea:





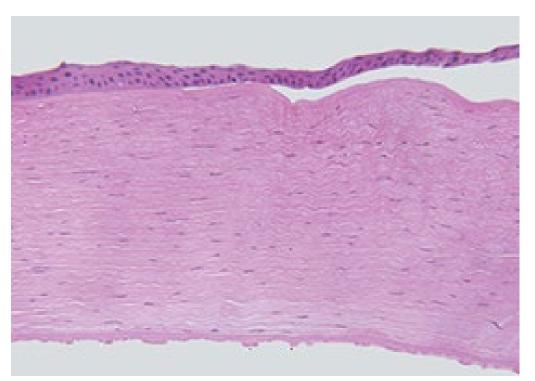
What's going on here?

- --The epi looks bad—it's thin in places, and over half has separated from Bowman's.
- --Speaking of: Bowman's seems intact
- --The stroma isn't taking a special stain, suggesting this isn't a stromal dystrophy.
- --Speaking of: Stromal clefting is almost absent, suggesting scarring and/or edema is present Turning our attention to the posterior cornea:
- --Descemet's appears a bit thir

thinned vs thickened

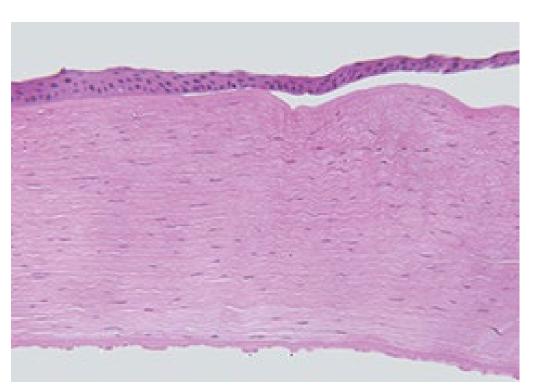
(admittedly this is a tough call to make)





- --The epi looks bad—it's thin in places, and over half has separated from Bowman's.
- --Speaking of: Bowman's seems intact
- --The stroma isn't taking a special stain, suggesting this isn't a stromal dystrophy.
- --Speaking of: Stromal clefting is almost absent, suggesting scarring and/or edema is present Turning our attention to the posterior cornea:
- --Descemet's appears a bit thickened (admittedly this is a tough call to make)



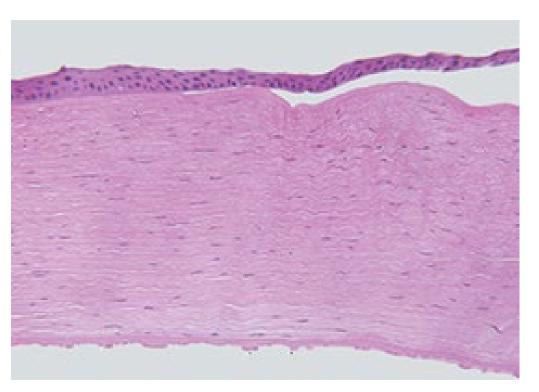


What's going on here?

- --The epi looks bad—it's thin in places, and over half has separated from Bowman's.
- --Speaking of: Bowman's seems intact
- --The stroma isn't taking a special stain, suggesting this isn't a stromal dystrophy.
- --Speaking of: Stromal clefting is almost absent, suggesting scarring and/or edema is present Turning our attention to the posterior cornea:
- --Descemet's appears a bit thickened (admittedly this is a tough call to make)
- --The endothelium is

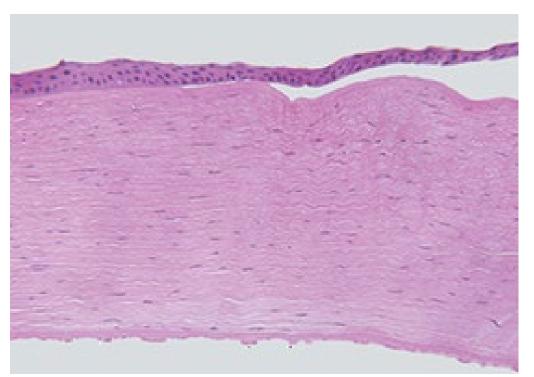
involved v uninvolved





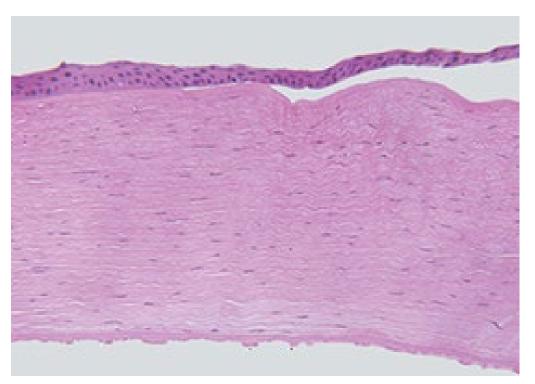
- --The epi looks bad—it's thin in places, and over half has separated from Bowman's.
- --Speaking of: Bowman's seems intact
- --The stroma isn't taking a special stain, suggesting this isn't a stromal dystrophy.
- --Speaking of: Stromal clefting is almost absent, suggesting scarring and/or edema is present Turning our attention to the posterior cornea:
- --Descemet's appears a bit thickened (admittedly this is a tough call to make)
- -- The endothelium is involved





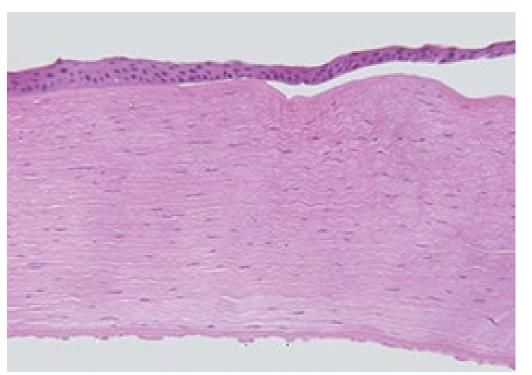
- --The epi looks bad—it's thin in places, and over half has separated from Bowman's.
- --Speaking of: Bowman's seems intact
- --The stroma isn't taking a special stain, suggesting this isn't a stromal dystrophy.
- --Speaking of: Stromal clefting is almost absent, suggesting scarring and/or edema is present Turning our attention to the posterior cornea:
- --Descemet's appears a bit thickened (admittedly this is a tough call to make)
- --The endothelium is involved as suggested by





- --The epi looks bad—it's thin in places, and over half has separated from Bowman's.
- --Speaking of: Bowman's seems intact
- --The stroma isn't taking a special stain, suggesting this isn't a stromal dystrophy.
- --Speaking of: Stromal clefting is almost absent, suggesting scarring and/or edema is present Turning our attention to the posterior cornea:
- --Descemet's appears a bit thickened (admittedly this is a tough call to make)
- --The endothelium is involved as suggested by the absence of endothelial cells.

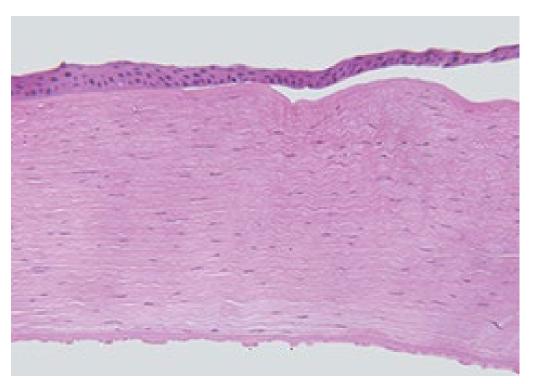




- --The epi looks bad—it's thin in places, and over half has separated from Bowman's.
- --Speaking of: Bowman's seems intact
- --The stroma isn't taking a special stain, suggesting this isn't a stromal dystrophy.
- --Speaking of: Stromal clefting is almost absent, suggesting scarring and/or edema is present Turning our attention to the posterior cornea:
- --Descemet's appears a bit thickened (admittedly this is a tough call to make)
- --The endothelium is involved as suggested by the absence of endothelial cells.

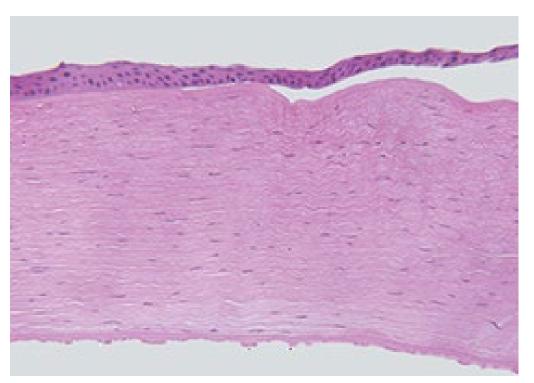
| -Descemet | several words | are | present |
|-----------|---------------|-----|---------|
| | | | |





- --The epi looks bad—it's thin in places, and over half has separated from Bowman's.
- --Speaking of: Bowman's seems intact
- --The stroma isn't taking a special stain, suggesting this isn't a stromal dystrophy.
- --Speaking of: Stromal clefting is almost absent, suggesting scarring and/or edema is present Turning our attention to the posterior cornea:
- --Descemet's appears a bit thickened (admittedly this is a tough call to make)
- --The endothelium is involved as suggested by the absence of endothelial cells.
- --Descemet excrescences (guttae) are present





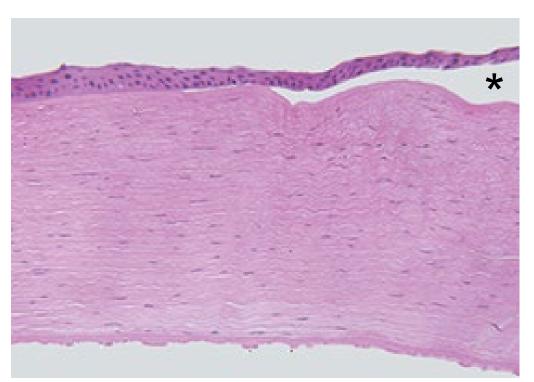
What's the diagnosis?

What's going on here?

- --The epi looks bad—it's thin in places, and over half has separated from Bowman's.
- --Speaking of: Bowman's seems intact
- --The stroma isn't taking a special stain, suggesting this isn't a stromal dystrophy.
- --Speaking of: Stromal clefting is almost absent, suggesting scarring and/or edema is present Turning our attention to the posterior cornea:
- --Descemet's appears a bit thickened (admittedly this is a tough call to make)
- --The endothelium is involved as suggested by the absence of endothelial cells.
- --Descemet excrescences (guttae) are present

Guttata + absent endo cells + stromal changes c/w edema + epi changes c/w edema points toward one dx:





What's the diagnosis?

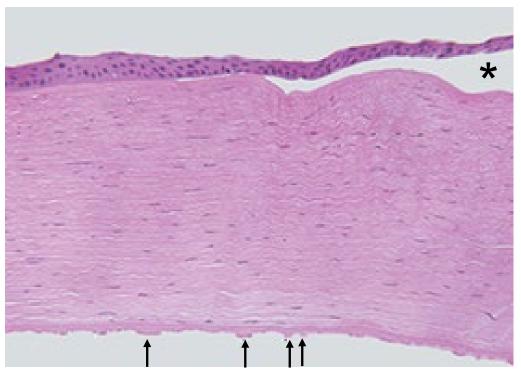
What's going on here?

- --The epi looks bad—it's thin in places, and over half has separated from Bowman's.
- --Speaking of: Bowman's seems intact
- --The stroma isn't taking a special stain, suggesting this isn't a stromal dystrophy.
- --Speaking of: Stromal clefting is almost absent, suggesting scarring and/or edema is present Turning our attention to the posterior cornea:
- --Descemet's appears a bit thickened (admittedly this is a tough call to make)
- --The endothelium is involved as suggested by the absence of endothelial cells.
- --Descemet excrescences (guttae) are present

Guttata + absent endo cells + stromal changes c/w edema + epi changes c/w edema points toward one dx:

Fuchs endothelial corneal dystrophy (FECD) is characterized by corneal edema, which can be severe enough to produce epithelial bullae (as in this example, *asterisk*).





What's the diagnosis?

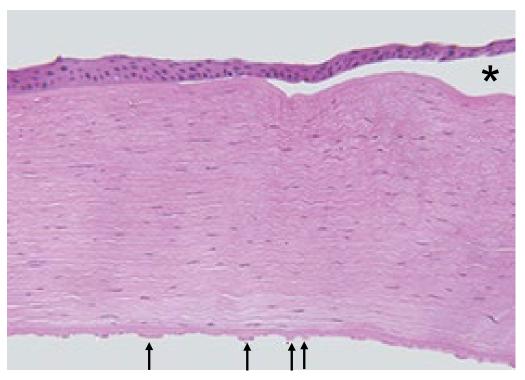
What's going on here?

- --The epi looks bad—it's thin in places, and over half has separated from Bowman's.
- --Speaking of: Bowman's seems intact
- --The stroma isn't taking a special stain, suggesting this isn't a stromal dystrophy.
- --Speaking of: Stromal clefting is almost absent, suggesting scarring and/or edema is present Turning our attention to the posterior cornea:
- --Descemet's appears a bit thickened (admittedly this is a tough call to make)
- --The endothelium is involved as suggested by the absence of endothelial cells.
- --Descemet excrescences (guttae) are present

Guttata + absent endo cells + stromal changes c/w edema + epi changes c/w edema points toward one dx:

Fuchs endothelial corneal dystrophy (FECD) is characterized by corneal edema, which can be severe enough to produce epithelial bullae (as in this example, *asterisk*). Descemet's becomes irregularly thickened, and guttae develop on it (*arrows*).





What's the diagnosis?

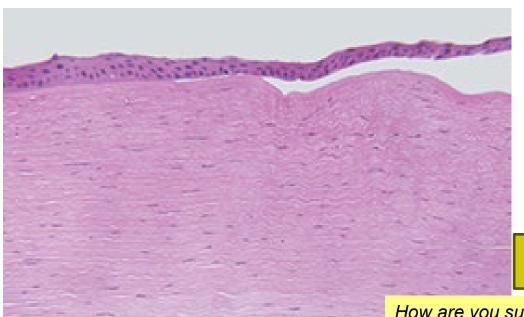
What's going on here?

- --The epi looks bad—it's thin in places, and over half has separated from Bowman's.
- --Speaking of: Bowman's seems intact
- --The stroma isn't taking a special stain, suggesting this isn't a stromal dystrophy.
- --Speaking of: Stromal clefting is almost absent, suggesting scarring and/or edema is present Turning our attention to the posterior cornea:
- --Descemet's appears a bit thickened (admittedly this is a tough call to make)
- --The endothelium is involved as suggested by the absence of endothelial cells.
- --Descemet excrescences (guttae) are present

Guttata + absent endo cells + stromal changes c/w edema + epi changes c/w edema points toward one dx:

Fuchs endothelial corneal dystrophy (FECD) is characterized by corneal edema, which can be severe enough to produce epithelial bullae (as in this example, *asterisk*). Descemet's becomes irregularly thickened, and guttae develop on it (*arrows*). The underlying pathology is loss of endothelial cell function and viability.





What's going on here?

- --The epi looks bad—it's thin in places, and over half has separated from Bowman's.
- --Speaking of: Bowman's seems intact
- --The stroma isn't taking a special stain, suggesting this isn't a stromal dystrophy.
- --Speaking of: Stromal clefting is almost absent, suggesting scarring and/or edema is present Turning our attention to the posterior cornea:
- --Descemet's appears a bit thickened (admittedly this is a tough call to make)

the absence of endothelial cells.

ggested by

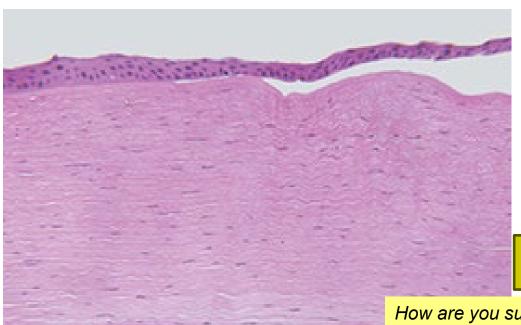
are present

How are you supposed to know the endothelial cell count is low?

What's the diagnosis?

Fuchs endothelial corneal dystrophy (I severe enough to produce epithelial bulla





What's going on here?

- --The epi looks bad—it's thin in places, and over half has separated from Bowman's.
- --Speaking of: Bowman's seems intact
- --The stroma isn't taking a special stain, suggesting this isn't a stromal dystrophy.
- --Speaking of: Stromal clefting is almost absent, suggesting scarring and/or edema is present Turning our attention to the posterior cornea:
- --Descemet's appears a bit thickened (admittedly this is a tough call to make)

the absence of endothelial cells.

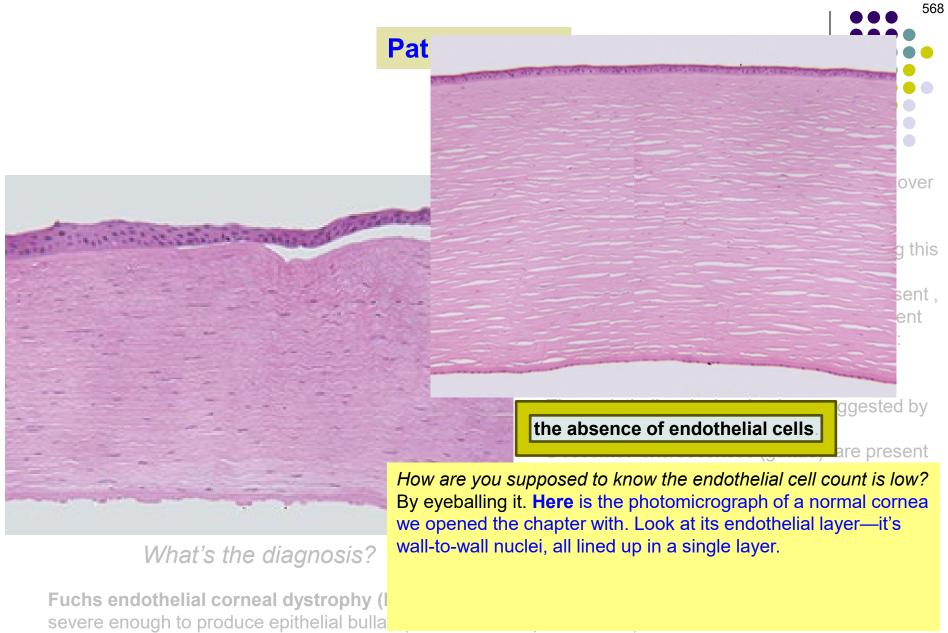
ggested by

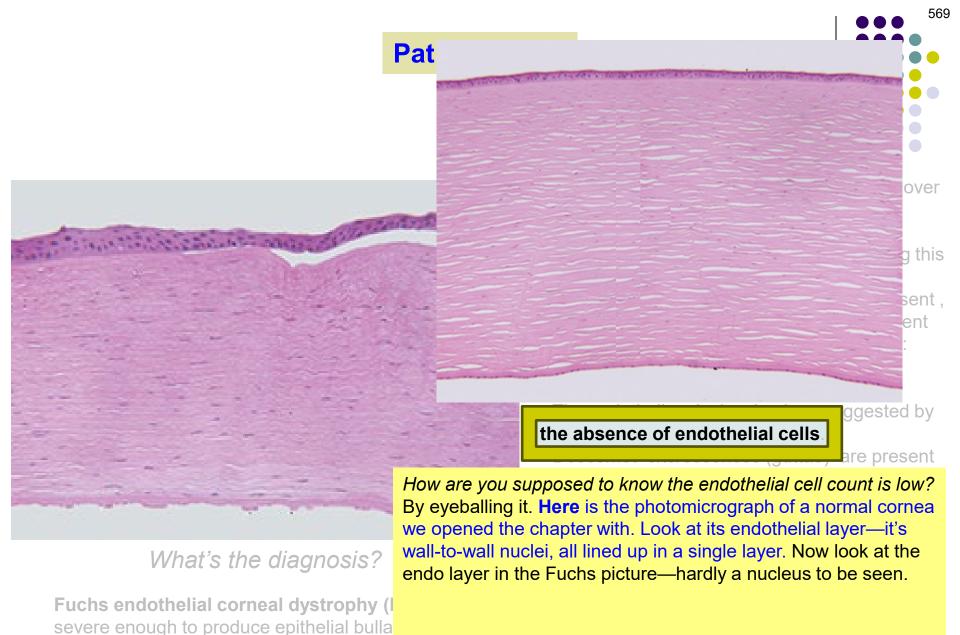
are present

How are you supposed to know the endothelial cell count is low? By eyeballing it

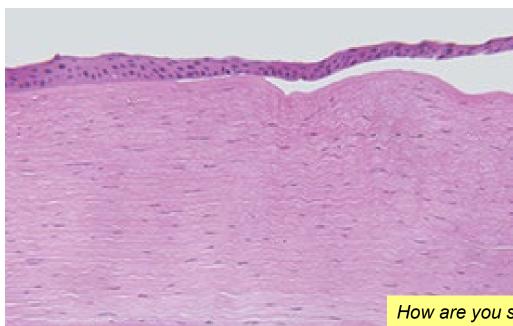
What's the diagnosis?

Fuchs endothelial corneal dystrophy (I severe enough to produce epithelial bulla









What's going on here?

- --The epi looks bad—it's thin in places, and over half has separated from Bowman's.
- --Speaking of: Bowman's seems intact
- --The stroma isn't taking a special stain, suggesting this isn't a stromal dystrophy.
- --Speaking of: Stromal clefting is almost absent, suggesting scarring and/or edema is present Turning our attention to the posterior cornea:
- --Descemet's appears a bit thickened (admittedly this is a tough call to make)

the absence of endothelial cells

are present

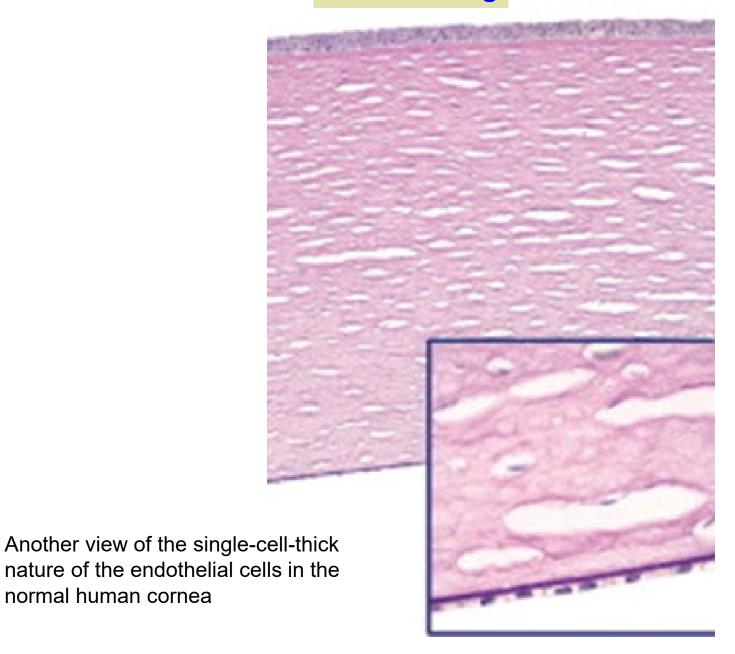
ggested by

How are you supposed to know the endothelial cell count is low? By eyeballing it. **Here** is the photomicrograph of a normal cornea we opened the chapter with. Look at its endothelial layer—it's wall-to-wall nuclei, all lined up in a single layer. Now look at the endo layer in the Fuchs picture—hardly a nucleus to be seen. If a single layer of tightly-spaced nuclei isn't found in an image, there's a problem with the endothelium.

What's the diagnosis?

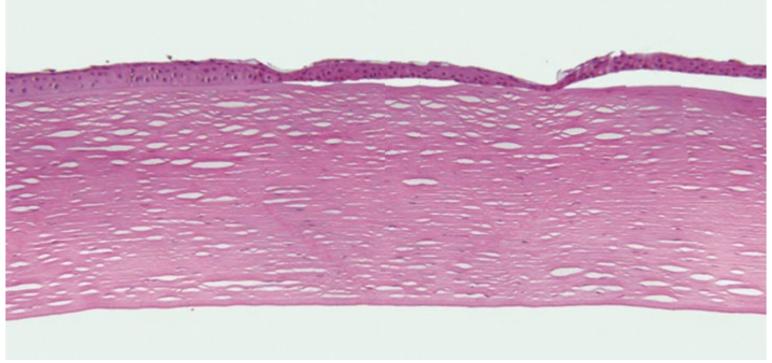
Fuchs endothelial corneal dystrophy (I severe enough to produce epithelial bulla

normal human cornea





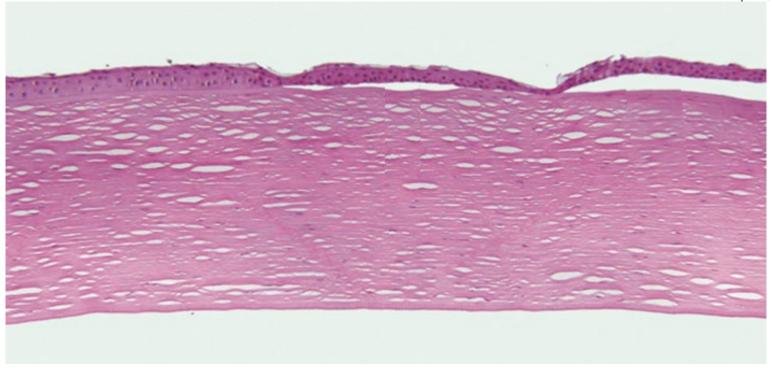




Here we have what seems like another case of FECD, given the:

- --?
- --?
- --?





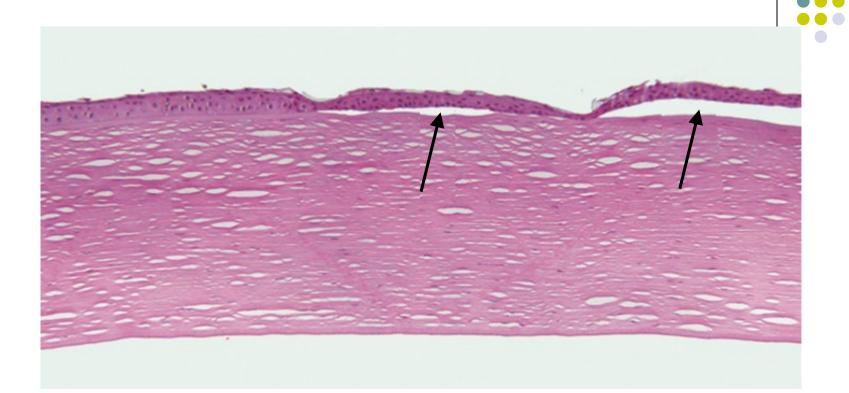
Here we have what seems like another case of FECD, given the:

--Epithelial

--?

--?

574



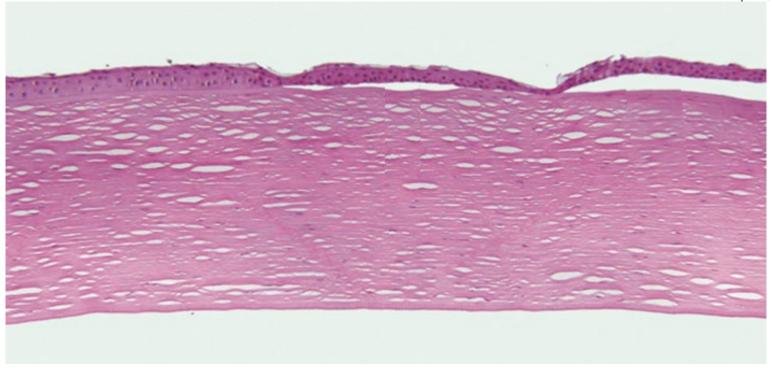
Here we have what seems like another case of FECD, given the:

--Epithelial bullae

--?

-- 2





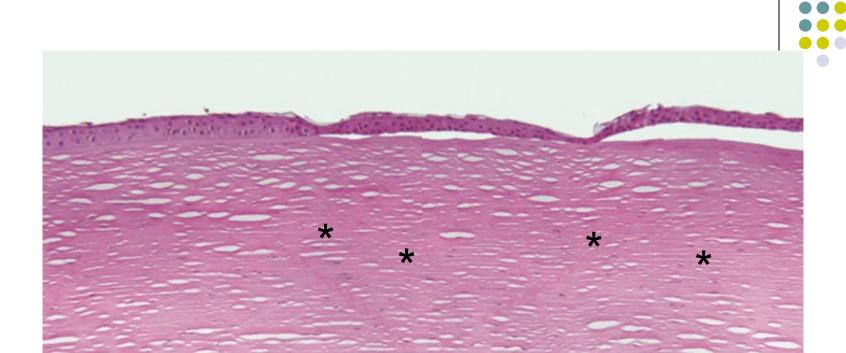
Here we have what seems like another case of FECD, given the:

--Epithelial bullae

--Loss of stromal c/w

--?

576

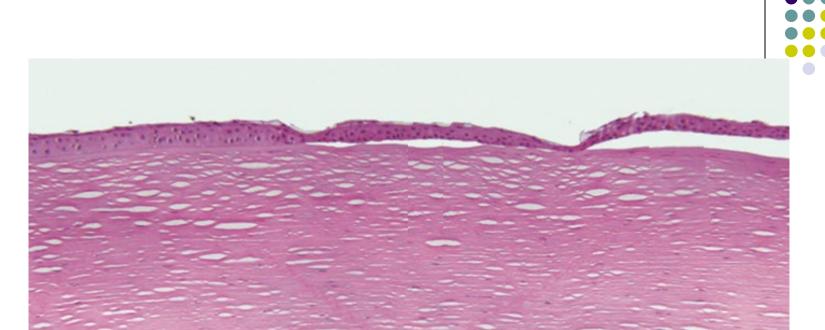


Here we have what seems like another case of FECD, given the:

- --Epithelial bullae
- --Loss of stromal clefting c/w edema

--?

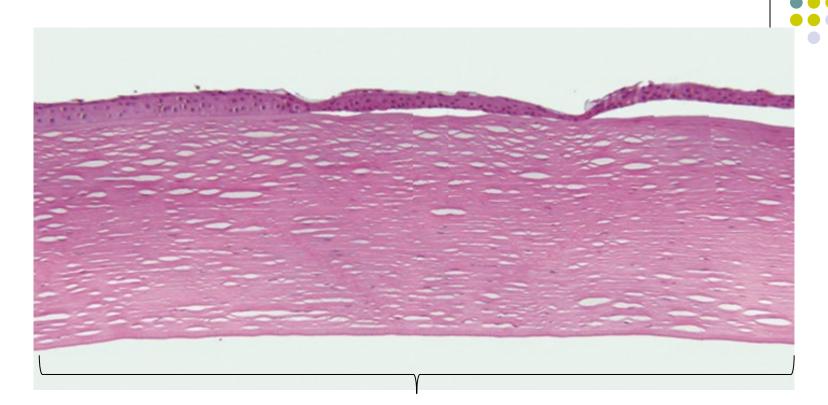
577



Here we have what seems like another case of FECD, given the:

- --Epithelial bullae
- --Loss of stromal clefting c/w edema
- --The absence of cells

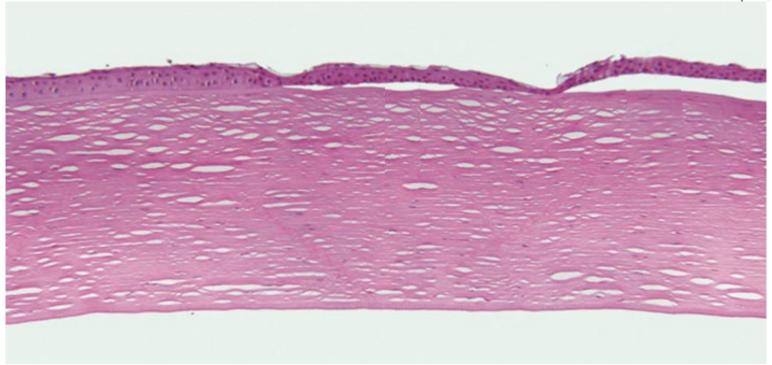
578



Here we have what seems like another case of FECD, given the:

- --Epithelial bullae
- --Loss of stromal clefting c/w edema
- -- The absence of endothelial cells





Here we have what seems like another case of FECD, given the:

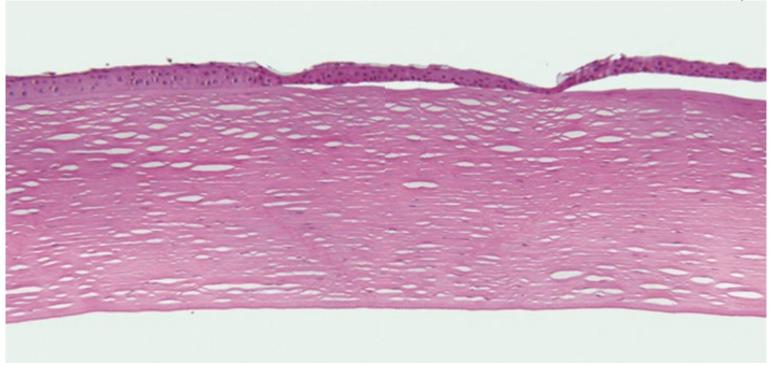
- --Epithelial bullae
- --Loss of stromal clefting c/w edema
- -- The absence of endothelial cells

But it mos def isn't FECD. How can you tell?

--?

--?





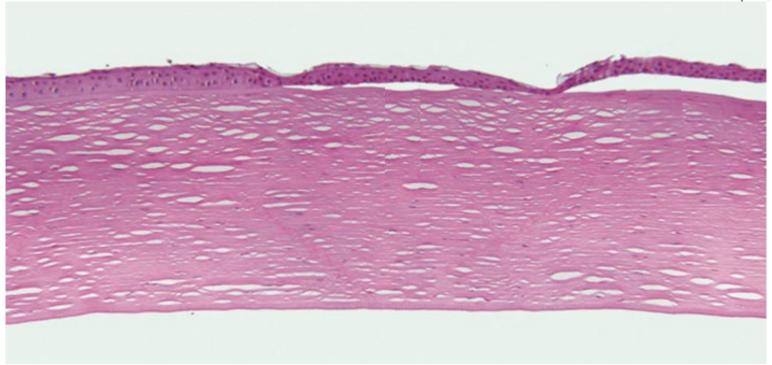
Here we have what seems like another case of FECD, given the:

- --Epithelial bullae
- --Loss of stromal clefting c/w edema
- -- The absence of endothelial cells

But it mos def isn't FECD. How can you tell?

--Descemet's isn't thickened vs thinned





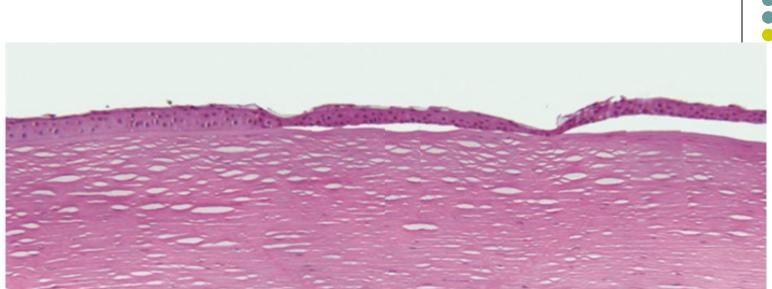
Here we have what seems like another case of FECD, given the:

- --Epithelial bullae
- --Loss of stromal clefting c/w edema
- -- The absence of endothelial cells

But it mos def isn't FECD. How can you tell?

--Descemet's isn't thickened (again, a tough call)

-- 7

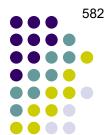


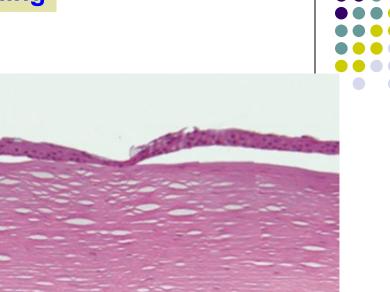
Here we have what seems like another case of FECD, given the:

- --Epithelial bullae
- --Loss of stromal clefting c/w edema
- -- The absence of endothelial cells

But it mos def isn't FECD. How can you tell?

- --Descemet's isn't thickened (again, a tough call)
- --The absence of





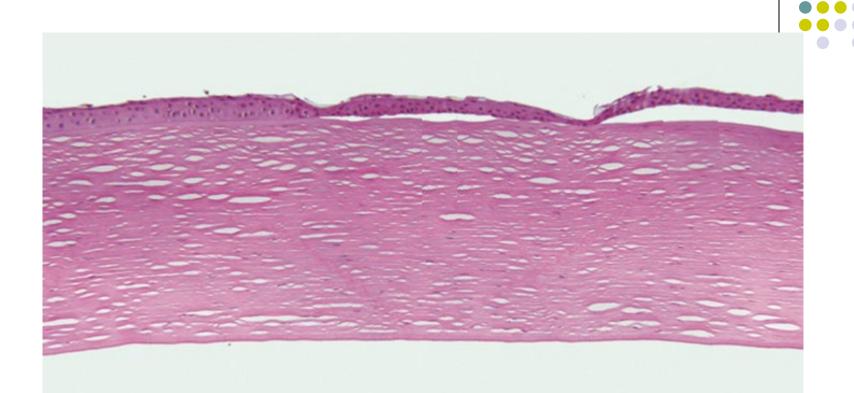
583

Here we have what seems like another case of FECD, given the:

- --Epithelial bullae
- --Loss of stromal clefting c/w edema
- -- The absence of endothelial cells

But it mos def isn't FECD. How can you tell?

- --Descemet's isn't thickened (again, a tough call)
- --The absence of guttata

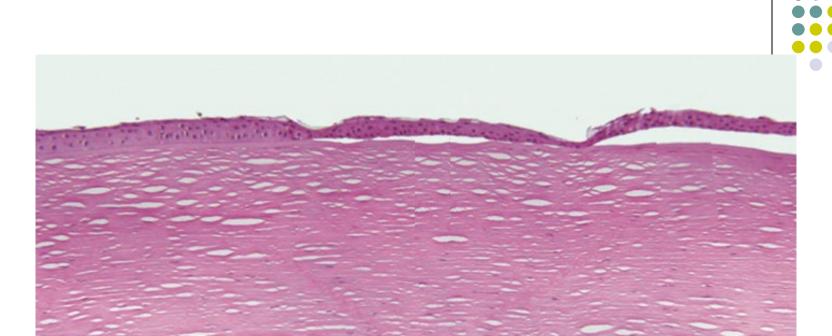


Here we have what seems like another case of FECD, given the:

- --Epithelial bullae
- --Loss of stromal clefting c/w edema
- --The absence of endothelial cells But it mos def *isn't* FECD. *How can you tell?*
- --Descemet's isn't thickened (again, a tough call)
- -- The absence of guttata

So, this is a case of two words 2ndry to endothelial-cell loss, but it's not FECD.

584



Here we have what seems like another case of FECD, given the:

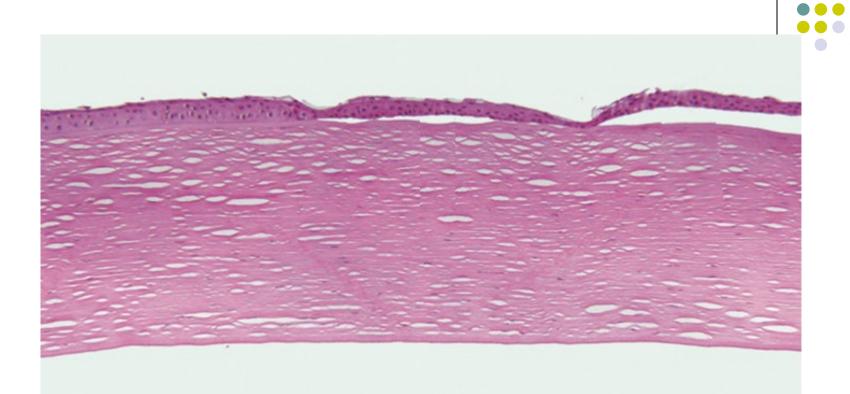
- --Epithelial bullae
- --Loss of stromal clefting c/w edema
- -- The absence of endothelial cells

But it mos def isn't FECD. How can you tell?

- --Descemet's isn't thickened (again, a tough call)
- -- The absence of guttata

So, this is a case of bullous keratopathy 2ndry to endothelial-cell loss, but it's not FECD.

585



Here we have what seems like another case of FECD, given the:

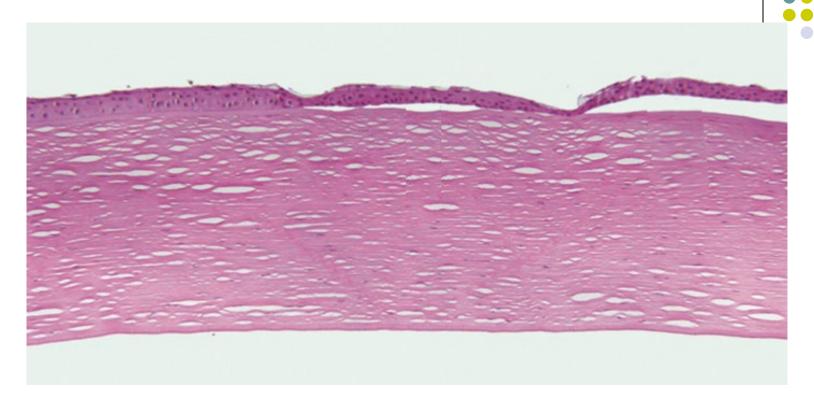
- --Epithelial bullae
- --Loss of stromal clefting c/w edema
- --The absence of endothelial cells

But it mos def isn't FECD. How can you tell?

- --Descemet's isn't thickened (again, a tough call)
- -- The absence of guttata

So, this is a case of bullous keratopathy 2ndry to endothelial-cell loss, but it's not FECD. What then is the most likely cause?

586



Here we have what seems like another case of FECD, given the:

- --Epithelial bullae
- --Loss of stromal clefting c/w edema
- --The absence of endothelial cells

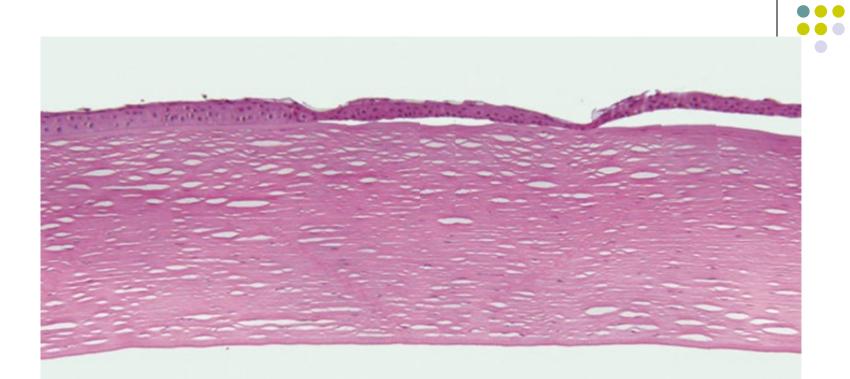
But it mos def isn't FECD. How can you tell?

- --Descemet's isn't thickened (again, a tough call)
- --The absence of guttata

So, this is a case of bullous keratopathy 2ndry to endothelial-cell loss, but it's not FECD. What then is the most likely cause?

587

Pseudophakic bullous keratopathy (PBK), which is in fact what's going on here.



Here we have what seems like another case of FECD, given the:

- --Epithelial bullae
- --Loss of stromal clefting c/w edema
- -- The absence of endothelial cells

But it mos def isn't FECD. How can you tell?

- --Descemet's isn't thickened (again, a tough call)
- --The absence of guttata

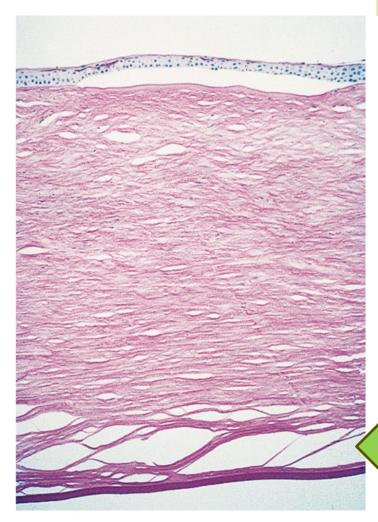
So, this is a case of bullous keratopathy 2ndry to endothelial-cell loss, but it's not FECD. What then is the most likely cause?

588

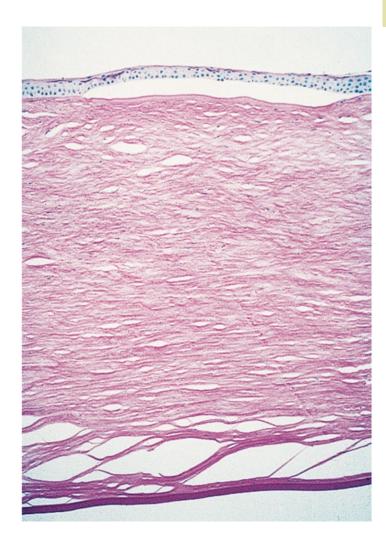
Pseudophakic bullous keratopathy (PBK), which is in fact what's going on here. The key to differentiating PBK from FECD is in noting the presence vs absence of guttata.







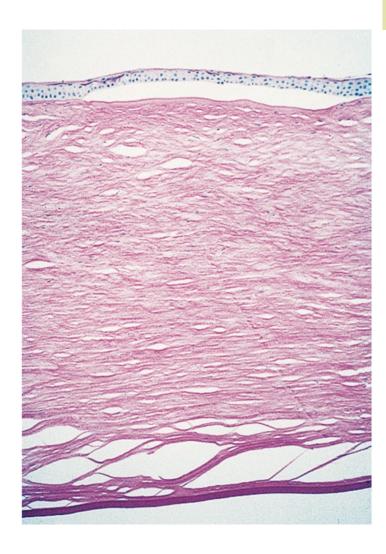
Before we get started, *note*: All this clefting is artifactual (and not the good kind we've been relying on). Just ignore it.





Here we have what seems like another case of PBK, as we have:

- --?
- --?
- --?
- --?





Here we have what seems like another case of PBK, as we have:

- --Epithelial bullae
- --Loss of stromal clefting c/w edema
- -- The absence of endothelial cells
- --No guttata





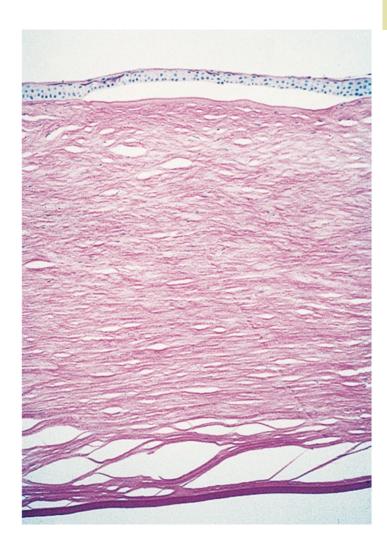
Here we have what seems like another case of PBK, as we have:

- --Epithelial bullae
- --Loss of stromal clefting c/w edema
- -- The absence of endothelial cells
- --No guttata

But it *isn't* PBK; rather, it is

words 1-3 of 4

word 4

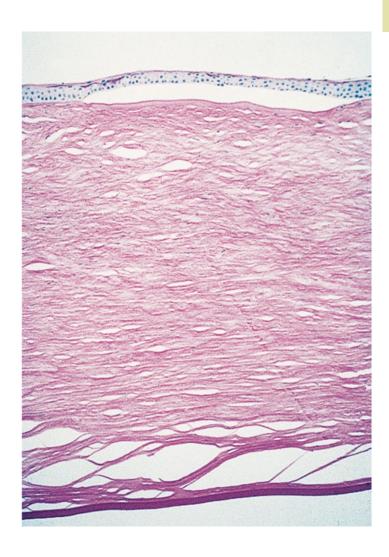




Here we have what seems like another case of PBK, as we have:

- --Epithelial bullae
- --Loss of stromal clefting c/w edema
- -- The absence of endothelial cells
- --No guttata

But it *isn't* PBK; rather, it is **congenital hereditary endothelial dystrophy** (CHED).



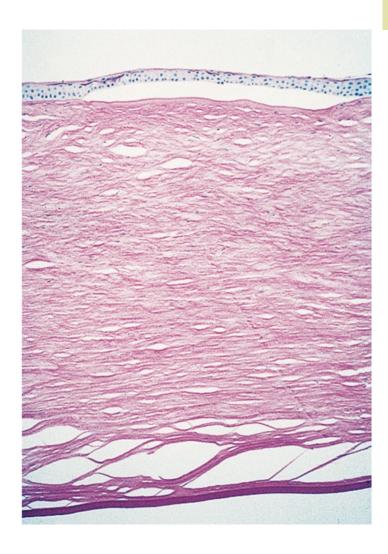


Here we have what seems like another case of PBK, as we have:

- --Epithelial bullae
- --Loss of stromal clefting c/w edema
- -- The absence of endothelial cells
- --No guttata

But it *isn't* PBK; rather, it is **congenital hereditary endothelial dystrophy** (CHED).

How can you tell it's CHED and not PBK?





Here we have what seems like another case of PBK, as we have:

- --Epithelial bullae
- --Loss of stromal clefting c/w edema
- -- The absence of endothelial cells
- --No guttata

But it *isn't* PBK; rather, it is **congenital hereditary endothelial dystrophy** (CHED).

How can you tell it's CHED and not PBK?

Because unlike in PBK, in CHED is thickened.



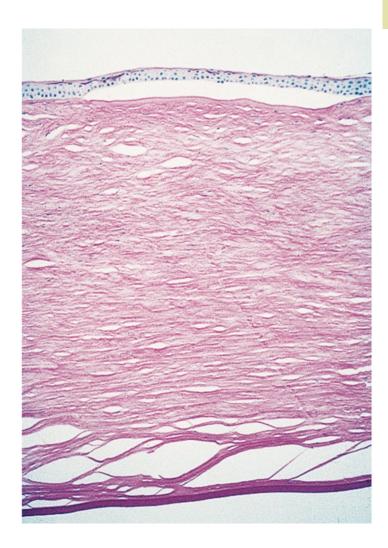


Here we have what seems like another case of PBK, as we have:

- --Epithelial bullae
- --Loss of stromal clefting c/w edema
- -- The absence of endothelial cells
- --No guttata

But it *isn't* PBK; rather, it is **congenital hereditary endothelial dystrophy** (CHED).

How can you tell it's CHED and not PBK?
Because unlike in PBK, in CHED Descemet's is thickened.





Here we have what seems like another case of PBK, as we have:

- --Epithelial bullae
- --Loss of stromal clefting c/w edema
- -- The absence of endothelial cells
- --No guttata

But it *isn't* PBK; rather, it is **congenital hereditary endothelial dystrophy** (CHED).

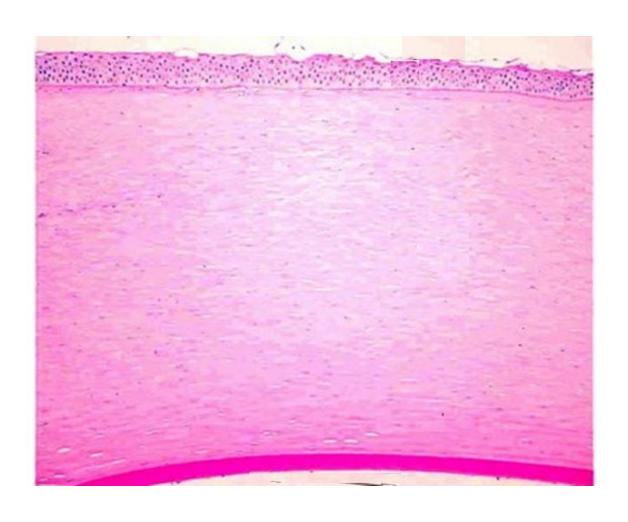
How can you tell it's CHED and not PBK?

Because unlike in PBK, in CHED Descemet's is thickened.

Again, tough call for those of us who aren't pathologists.

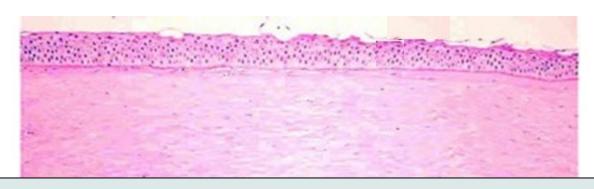
Should you be expected to make this call on a test, I reckon the thickening would be much more obvious than in this pic (eg, see the next slide).





CHED. Even I can tell Descemet's is thick here.



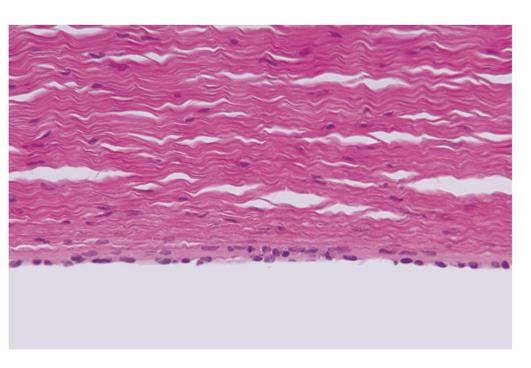


For more on FECD and CHED, see slide-set K45



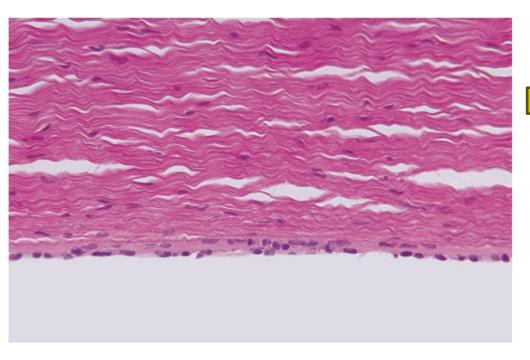
CHED. Even I can tell Descemet's is thick here.





It should not surprise that the key finding in **this** condition is to be found in the posterior cornea...

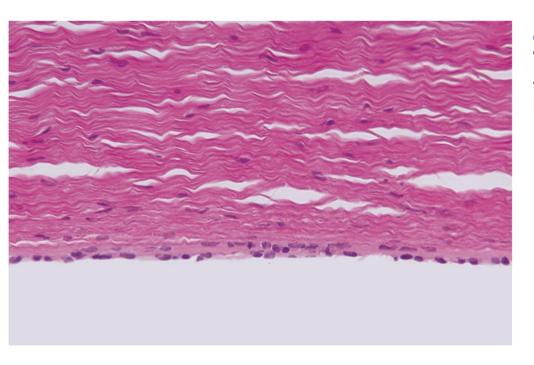




It should not surprise that the key finding in **this** condition is to be found in the posterior cornea...

What's going on here?
--The stroma is Y taking a stain, suggesting this is v isn't a stromal dystrophy.

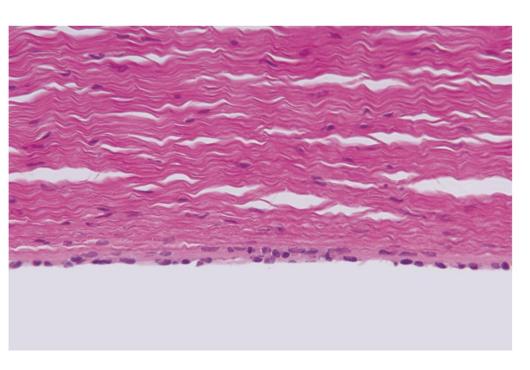




It should not surprise that the key finding in **this** condition is to be found in the posterior cornea... What's going on here?

--The stroma isn't taking a stain, suggesting this isn't a stromal dystrophy.





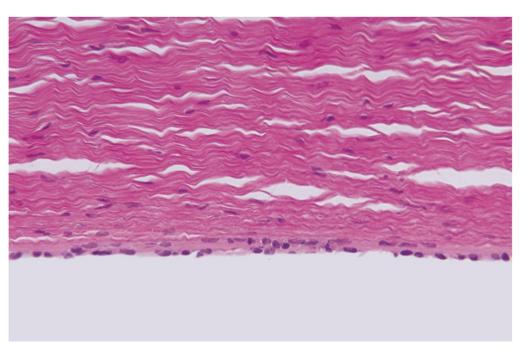
It should not surprise that the key finding in **this** condition is to be found in the posterior cornea...

What's going on here?

- --The stroma isn't taking a stain, suggesting this isn't a stromal dystrophy.
- --Speaking of: Stromal clefting is





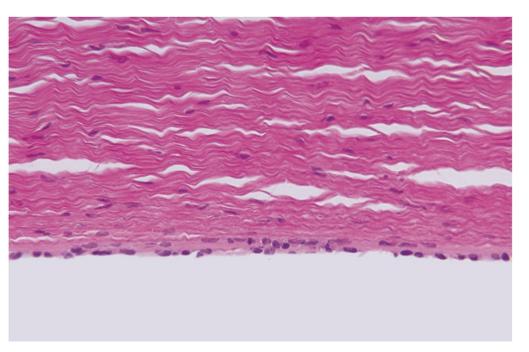


It should not surprise that the key finding in **this** condition is to be found in the posterior cornea...

What's going on here?

- --The stroma isn't taking a stain, suggesting this isn't a stromal dystrophy.
- --Speaking of: Stromal clefting is unaffected

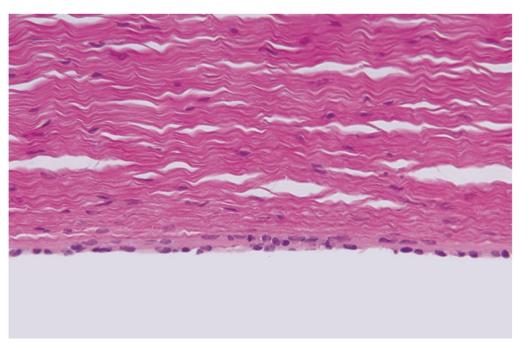




It should not surprise that the key finding in **this** condition is to be found in the posterior cornea... What's going on here?

- --The stroma isn't taking a stain, suggesting this isn't a stromal dystrophy.
- --Speaking of: Stromal clefting is unaffected Turning our attention to the posterior cornea:
- --Descemet's appears , for lack of a better term (I know—tough call again).

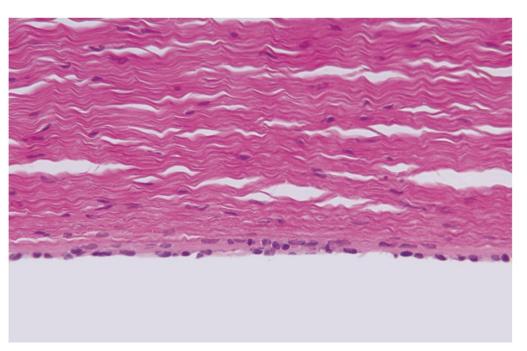




It should not surprise that the key finding in **this** condition is to be found in the posterior cornea... What's going on here?

- --The stroma isn't taking a stain, suggesting this isn't a stromal dystrophy.
- --Speaking of: Stromal clefting is unaffected Turning our attention to the posterior cornea:
- --Descemet's appears wonky, for lack of a better term (I know—tough call again).



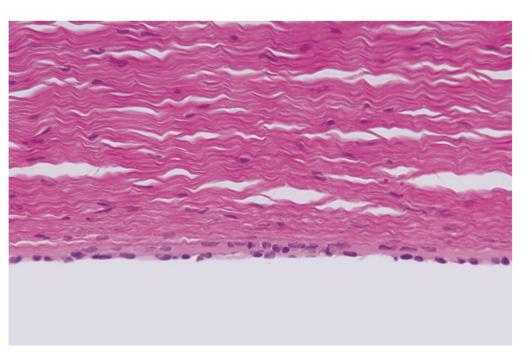


It should not surprise that the key finding in **this** condition is to be found in the posterior cornea... What's going on here?

- --The stroma isn't taking a stain, suggesting this isn't a stromal dystrophy.
- --Speaking of: Stromal clefting is unaffected Turning our attention to the posterior cornea:
- --Descemet's appears wonky, for lack of a better term (I know—tough call again).
- --OTOH, what's **not** a tough call is that the endothelium is affected vs unaffected as suggested by its

two words

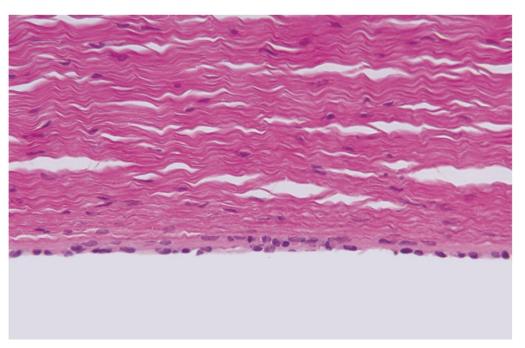




It should not surprise that the key finding in **this** condition is to be found in the posterior cornea... What's going on here?

- --The stroma isn't taking a stain, suggesting this isn't a stromal dystrophy.
- --Speaking of: Stromal clefting is unaffected Turning our attention to the posterior cornea:
- --Descemet's appears wonky, for lack of a better term (I know—tough call again).
- --OTOH, what's **not** a tough call is that the endothelium is jacked up as suggested by its multiple layers.





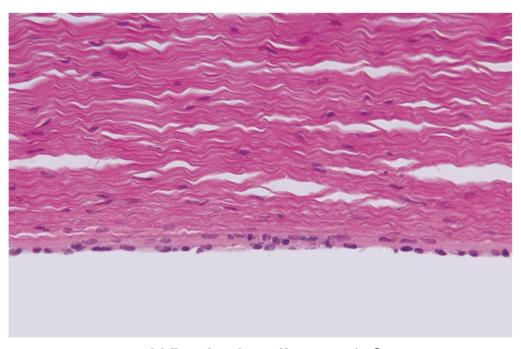
What's the diagnosis?

It should not surprise that the key finding in **this** condition is to be found in the posterior cornea... What's going on here?

- --The stroma isn't taking a stain, suggesting this isn't a stromal dystrophy.
- --Speaking of: Stromal clefting is unaffected Turning our attention to the posterior cornea:
- --Descemet's appears wonky, for lack of a better term (I know—tough call again).
- --OTOH, what's **not** a tough call is that the endothelium is jacked up as suggested by its multiple layers.

Wonky Descemet's + multilayered endothelium strongly indicates one dx:





It should not surprise that the key finding in **this** condition is to be found in the posterior cornea... What's going on here?

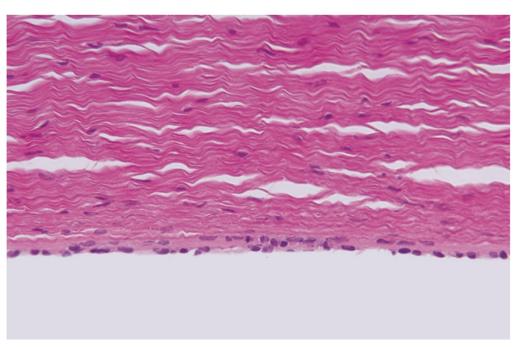
- --The stroma isn't taking a stain, suggesting this isn't a stromal dystrophy.
- --Speaking of: Stromal clefting is unaffected Turning our attention to the posterior cornea:
- --Descemet's appears wonky , for lack of a better term (I know—tough call again).
- --OTOH, what's **not** a tough call is that the endothelium is jacked up as suggested by its multiple layers.

Wonky Descemet's + multilayered endothelium strongly indicates one dx:

What's the diagnosis?

Posterior polymorphous dystrophy (PPMD) has a variety of clinical manifestations.





It should not surprise that the key finding in **this** condition is to be found in the posterior cornea... What's going on here?

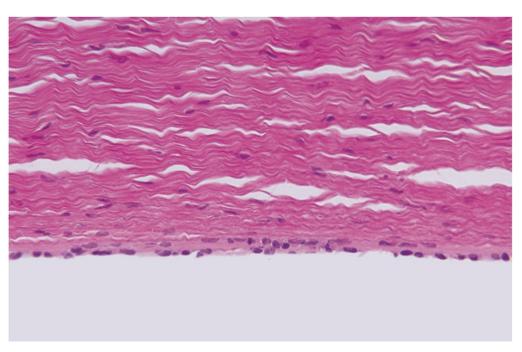
- --The stroma isn't taking a stain, suggesting this isn't a stromal dystrophy.
- --Speaking of: Stromal clefting is unaffected Turning our attention to the posterior cornea:
- --Descemet's appears wonky , for lack of a better term (I know—tough call again).
- --OTOH, what's **not** a tough call is that the endothelium is jacked up as suggested by its multiple layers.

Wonky Descemet's + multilayered endothelium strongly indicates one dx:

What's the diagnosis?

Posterior polymorphous dystrophy (PPMD) has a variety of clinical manifestations. Histologically, Descemet's is and





It should not surprise that the key finding in **this** condition is to be found in the posterior cornea... What's going on here?

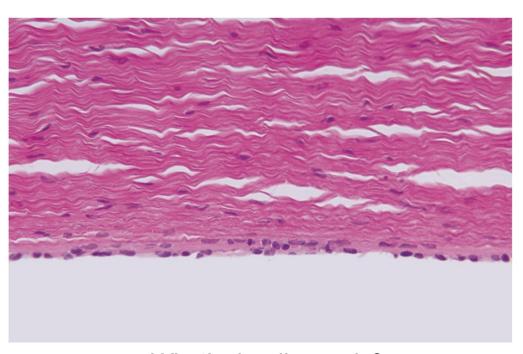
- --The stroma isn't taking a stain, suggesting this isn't a stromal dystrophy.
- --Speaking of: Stromal clefting is unaffected Turning our attention to the posterior cornea:
- --Descemet's appears wonky , for lack of a better term (I know—tough call again).
- --OTOH, what's **not** a tough call is that the endothelium is jacked up as suggested by its multiple layers.

Wonky Descemet's + multilayered endothelium strongly indicates one dx:

What's the diagnosis?

Posterior polymorphous dystrophy (PPMD) has a variety of clinical manifestations. Histologically, Descemet's is thickened and laminated





It should not surprise that the key finding in **this** condition is to be found in the posterior cornea... What's going on here?

- --The stroma isn't taking a stain, suggesting this isn't a stromal dystrophy.
- --Speaking of: Stromal clefting is unaffected Turning our attention to the posterior cornea:
- --Descemet's appears wonky , for lack of a better term (I know—tough call again).
- --OTOH, what's **not** a tough call is that the endothelium is jacked up as suggested by its multiple layers.

Wonky Descemet's + multilayered endothelium strongly indicates one dx:

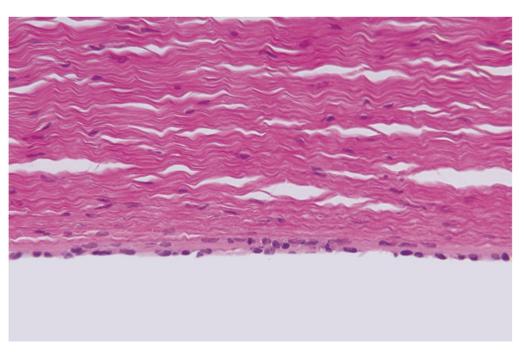
What's the diagnosis?

Posterior polymorphous dystrophy (PPMD) has a variety of clinical manifestations.

Histologically, Descemet's is thickened and laminated; excrescences

may be *v* are not present.





It should not surprise that the key finding in **this** condition is to be found in the posterior cornea... What's going on here?

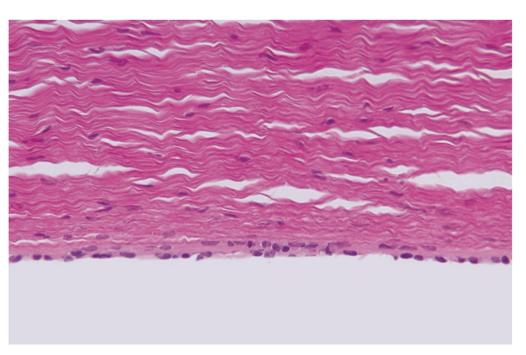
- --The stroma isn't taking a stain, suggesting this isn't a stromal dystrophy.
- --Speaking of: Stromal clefting is unaffected Turning our attention to the posterior cornea:
- --Descemet's appears wonky , for lack of a better term (I know—tough call again).
- --OTOH, what's **not** a tough call is that the endothelium is jacked up as suggested by its multiple layers.

Wonky Descemet's + multilayered endothelium strongly indicates one dx:

What's the diagnosis?

Posterior polymorphous dystrophy (PPMD) has a variety of clinical manifestations. Histologically, Descemet's is thickened and laminated; excrescences may be present.





It should not surprise that the key finding in **this** condition is to be found in the posterior cornea... What's going on here?

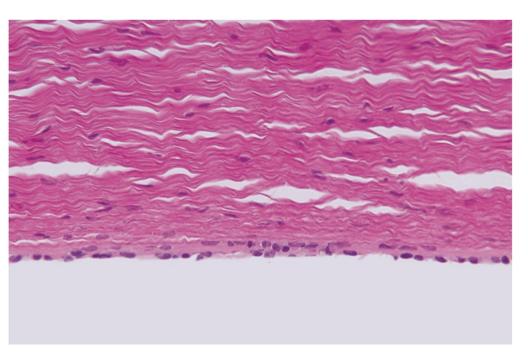
- --The stroma isn't taking a stain, suggesting this isn't a stromal dystrophy.
- --Speaking of: Stromal clefting is unaffected Turning our attention to the posterior cornea:
- --Descemet's appears wonky , for lack of a better term (I know—tough call again).
- --OTOH, what's **not** a tough call is that the endothelium is jacked up as suggested by its multiple layers.

Wonky Descemet's + multilayered endothelium strongly indicates one dx:

What's the diagnosis?

Posterior polymorphous dystrophy (PPMD) has a variety of clinical manifestations. Histologically, Descemet's is thickened and laminated; excrescences may be present. The underlying pathology is endothelial cell action





It should not surprise that the key finding in **this** condition is to be found in the posterior cornea... What's going on here?

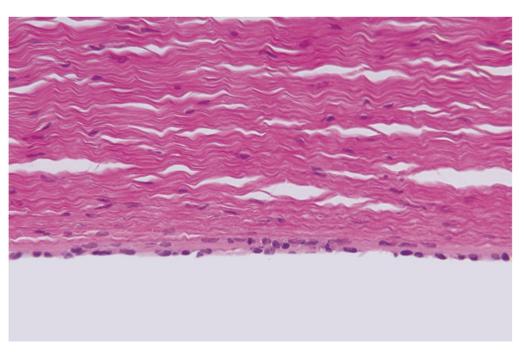
- --The stroma isn't taking a stain, suggesting this isn't a stromal dystrophy.
- --Speaking of: Stromal clefting is unaffected Turning our attention to the posterior cornea:
- --Descemet's appears wonky , for lack of a better term (I know—tough call again).
- --OTOH, what's **not** a tough call is that the endothelium is jacked up as suggested by its multiple layers.

Wonky Descemet's + multilayered endothelium strongly indicates one dx:

What's the diagnosis?

Posterior polymorphous dystrophy (PPMD) has a variety of clinical manifestations. Histologically, Descemet's is thickened and laminated; excrescences may be present. The underlying pathology is endothelial cell transformation





It should not surprise that the key finding in **this** condition is to be found in the posterior cornea... What's going on here?

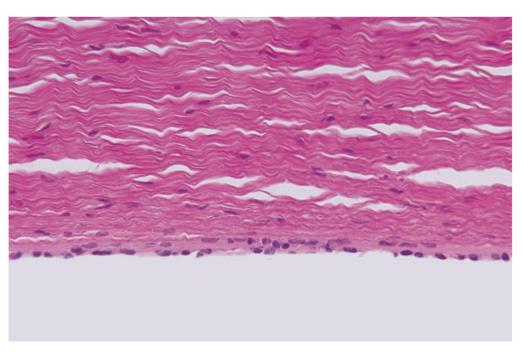
- --The stroma isn't taking a stain, suggesting this isn't a stromal dystrophy.
- --Speaking of: Stromal clefting is unaffected Turning our attention to the posterior cornea:
- --Descemet's appears wonky , for lack of a better term (I know—tough call again).
- --OTOH, what's **not** a tough call is that the endothelium is jacked up as suggested by its multiple layers.

Wonky Descemet's + multilayered endothelium strongly indicates one dx:

What's the diagnosis?

Posterior polymorphous dystrophy (PPMD) has a variety of clinical manifestations. Histologically, Descemet's is thickened and laminated; excrescences may be present. The underlying pathology is endothelial cell transformation, which leads to them looking and 'behaving' like cells and/or cell type





It should not surprise that the key finding in **this** condition is to be found in the posterior cornea... What's going on here?

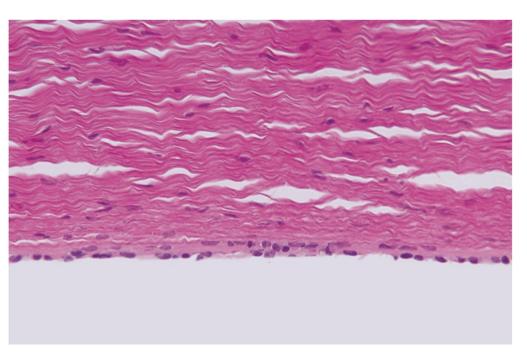
- --The stroma isn't taking a stain, suggesting this isn't a stromal dystrophy.
- --Speaking of: Stromal clefting is unaffected Turning our attention to the posterior cornea:
- --Descemet's appears wonky , for lack of a better term (I know—tough call again).
- --OTOH, what's **not** a tough call is that the endothelium is jacked up as suggested by its multiple layers.

Wonky Descemet's + multilayered endothelium strongly indicates one dx:

What's the diagnosis?

Posterior polymorphous dystrophy (PPMD) has a variety of clinical manifestations. Histologically, Descemet's is thickened and laminated; excrescences may be present. The underlying pathology is endothelial cell transformation, which leads to them looking and 'behaving' like epithelial cells and/or fibroblasts.





It should not surprise that the key finding in **this** condition is to be found in the posterior cornea... What's going on here?

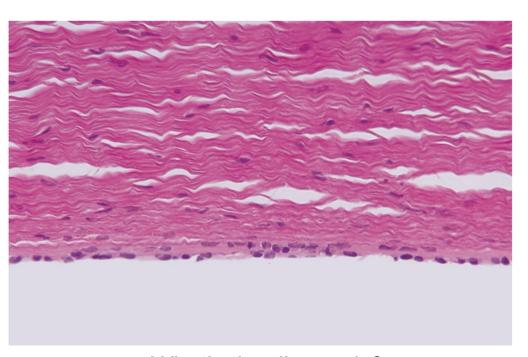
- --The stroma isn't taking a stain, suggesting this isn't a stromal dystrophy.
- --Speaking of: Stromal clefting is unaffected Turning our attention to the posterior cornea:
- --Descemet's appears wonky , for lack of a better term (I know—tough call again).
- --OTOH, what's **not** a tough call is that the endothelium is jacked up as suggested by its multiple layers.

Wonky Descemet's + multilayered endothelium strongly indicates one dx:

What's the diagnosis?

Posterior polymorphous dystrophy (PPMD) has a variety of clinical manifestations. Histologically, Descemet's is thickened and laminated; excrescences may be present. The underlying pathology is endothelial cell transformation, which leads to them looking and 'behaving' like epithelial cells and/or fibroblasts. (This explains how the endothelium comes to have its characteristic appearance.)





It should not surprise that the key finding in **this** condition is to be found in the posterior cornea... What's going on here?

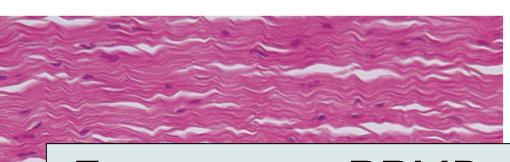
- --The stroma isn't taking a stain, suggesting this isn't a stromal dystrophy.
- --Speaking of: Stromal clefting is unaffected Turning our attention to the posterior cornea:
- --Descemet's appears wonky , for lack of a better term (I know—tough call again).
- --OTOH, what's **not** a tough call is that the endothelium is jacked up as suggested by its multiple layers.

Wonky Descemet's + multilayered endothelium strongly indicates one dx:

What's the diagnosis?

Posterior polymorphous dystrophy (PPMD) has a variety of clinical manifestations. Histologically, Descemet's is thickened and laminated; excrescences may be present. The underlying pathology is endothelial cell transformation, which leads to them looking and 'behaving' like epithelial cells and/or fibroblasts. (This explains how the endothelium comes to have its characteristic multilayered appearance.)





It should not surprise that the key finding in **this** condition is to be found in the posterior cornea... What's going on here?

- --The stroma isn't taking a stain, suggesting this isn't a stromal dystrophy.
- --Speaking of: Stromal clefting is unaffected

Turning our attention to the nectorior corner

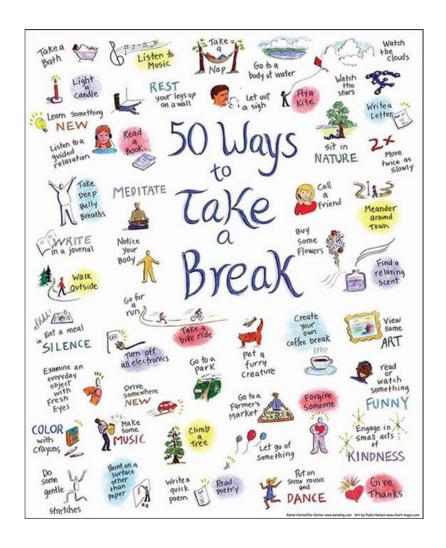
For more on PPMD, see slide-set K45

endothelium is jacked up as suggested by its multiple layers.

Wonky Descemet's + multilayered endothelium strongly indicates one dx:

What's the diagnosis?

Posterior polymorphous dystrophy (PPMD) has a variety of clinical manifestations. Histologically, Descemet's is thickened and laminated; excrescences may be present. The underlying pathology is endothelial cell transformation, which leads to them looking and 'behaving' like epithelial cells and/or fibroblasts. (This explains how the endothelium comes to have its characteristic multilayered appearance.)

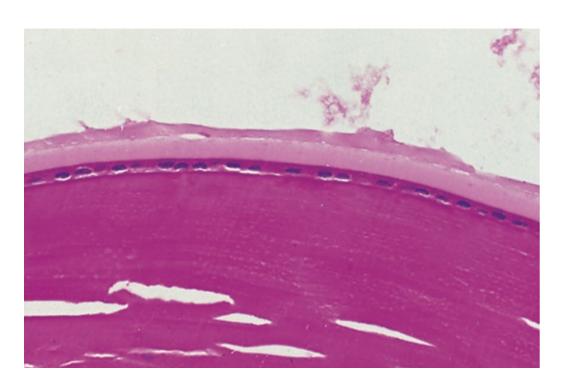


(This is a good point in the set to take a break)

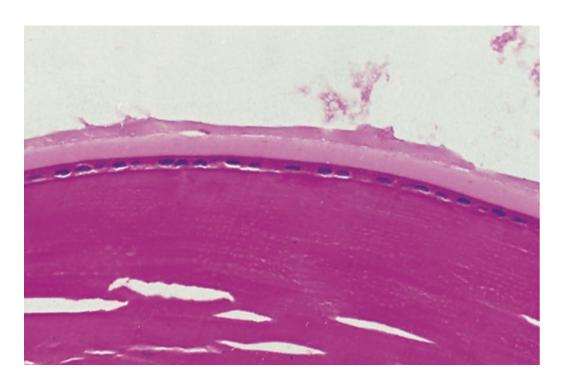




First: What tissue/structure is this?



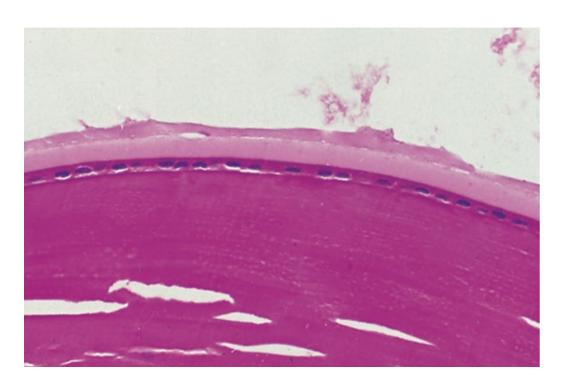




First: What tissue/structure is this?

It kinda looks like the anterior cornea—you can see what looks like stroma, a layer of epithelium, and Bowman's.



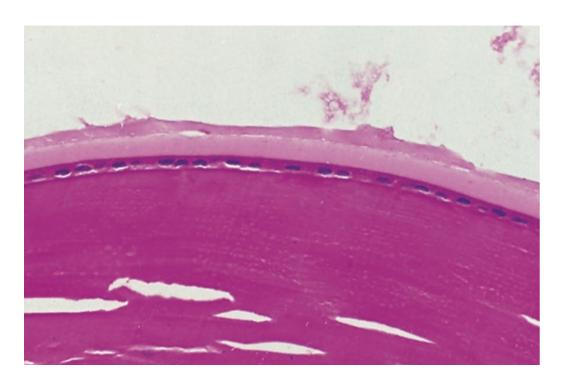


First: What tissue/structure is this?

It kinda looks like the anterior cornea—you can see what looks like stroma, a layer of epithelium, and Bowman's. But this interpretation is problematic in that:

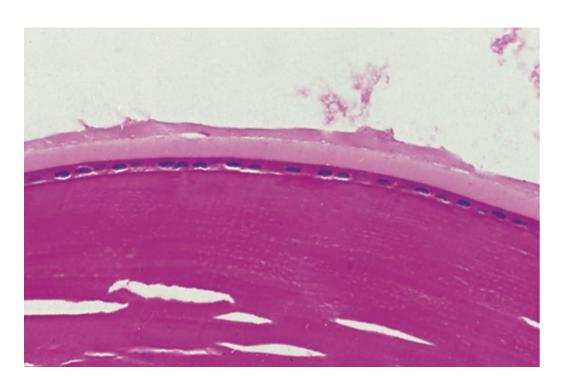
--It's a layered epi (K epi is layered)





First: What tissue/structure is this?
It kinda looks like the anterior cornea—you can see what looks like stroma, a layer of epithelium, and Bowman's. But this interpretation is problematic in that:
--It's a single-layered epi (K epi is multi-layered)



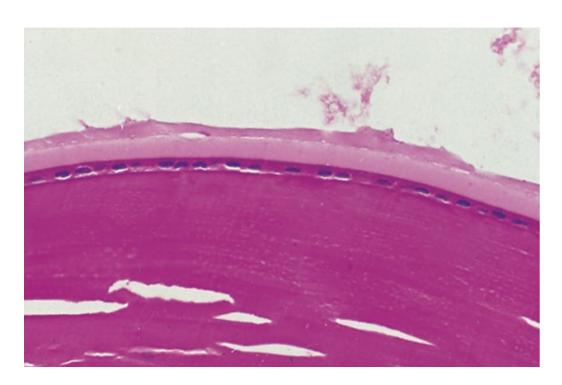


First: What tissue/structure is this?

It kinda looks like the anterior cornea—you can see what looks like stroma, a layer of epithelium, and Bowman's. But this interpretation is problematic in that:
--It's a single-layered epi (K epi is multi-layered)

--It's locale to the 'Bowman's' (K epi is locale of Bowman's)



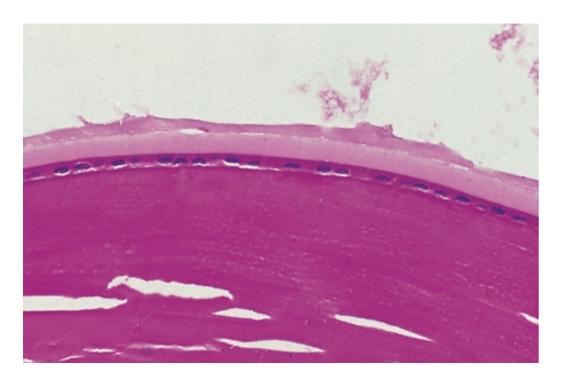


First: What tissue/structure is this?

It kinda looks like the anterior cornea—you can see what looks like stroma, a layer of epithelium, and Bowman's. But this interpretation is problematic in that:
--It's a single -layered epi (K epi is multi - layered)

--It's deep to the 'Bowman's' (K epi is on top of Bowman's)





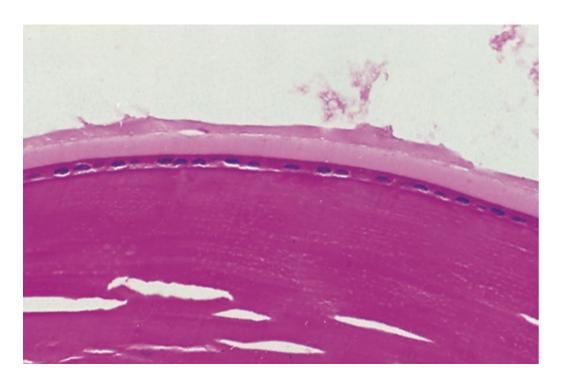
First: What tissue/structure is this?

It kinda looks like the anterior cornea—you can see what looks like stroma, a layer of epithelium, and Bowman's. But this interpretation is problematic in that:
--It's a single -layered epi (K epi is multi - layered)

--It's deep to the 'Bowman's' (K epi is on top of Bowman's)

So it ain't cornea.





First: What tissue/structure is this?

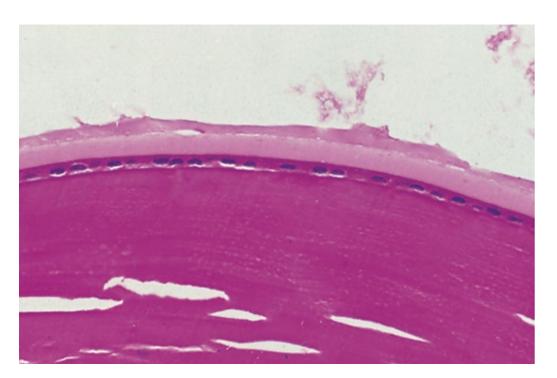
It kinda looks like the anterior cornea—you can see what looks like stroma, a layer of epithelium, and Bowman's. But this interpretation is problematic in that:
--It's a single -layered epi (K epi is multi - layered)

--It's deep to the 'Bowman's' (K epi is on top of Bowman's)

So it ain't cornea.

There's only one structure that has a membrane (with an epi *under* it) that surrounds amorphous material: ?





First: What tissue/structure is this?

It kinda looks like the anterior cornea—you can see what looks like stroma, a layer of epithelium, and Bowman's. But this interpretation is problematic in that:
--It's a single -layered epi (K epi is multi - layered)

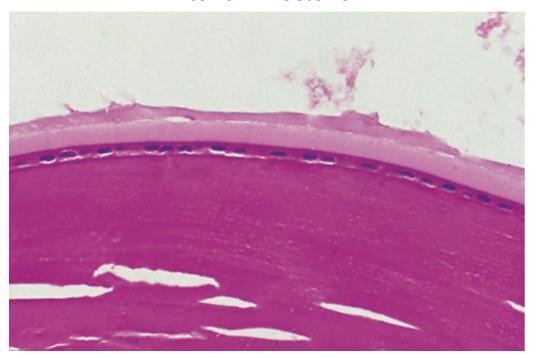
--It's deep to the 'Bowman's' (K epi is on top of Bowman's)

So it ain't cornea.

There's only one structure that has a membrane (with an epi *under* it) that surrounds amorphous material: The lens.



Anterior? Posterior?



First: What tissue/structure is this?

It kinda looks like the anterior cornea—you can see what looks like stroma, a layer of epithelium, and Bowman's. But this interpretation is problematic in that:
--It's a single -layered epi (K epi is multi - layered)

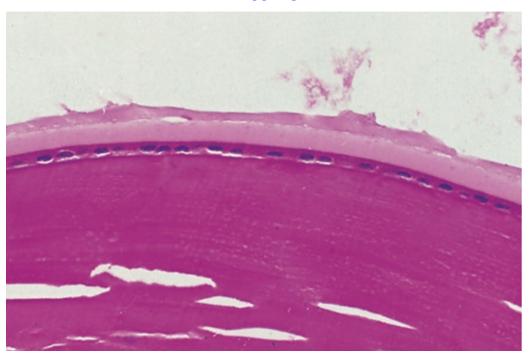
--It's deep to the 'Bowman's' (K epi is on top of Bowman's)

So it ain't cornea.

There's only one structure that has a membrane (with an epi *under* it) that surrounds amorphous material: The lens. *Is this the anterior or posterior lens?*



Anterior



First: What tissue/structure is this?

It kinda looks like the anterior cornea—you can see what looks like stroma, a layer of epithelium, and Bowman's. But this interpretation is problematic in that:
--It's a single -layered epi (K epi is multi - layered)

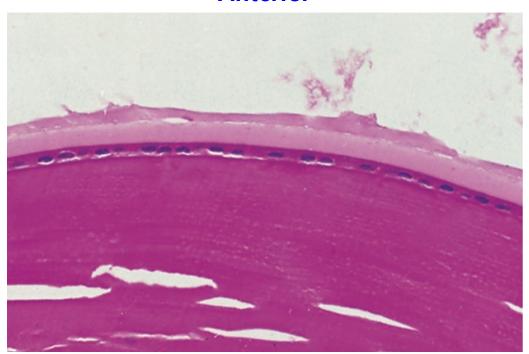
--It's deep to the 'Bowman's' (K epi is on top of Bowman's)

So it ain't cornea.

There's only one structure that has a membrane (with an epi *under* it) that surrounds amorphous material: The lens. *Is this the anterior or posterior lens?* Anterior.



Anterior



First: What tissue/structure is this?

It kinda looks like the anterior cornea—you can see what looks like stroma, a layer of epithelium, and Bowman's. But this interpretation is problematic in that:
--It's a single -layered epi (K epi is multi - layered)

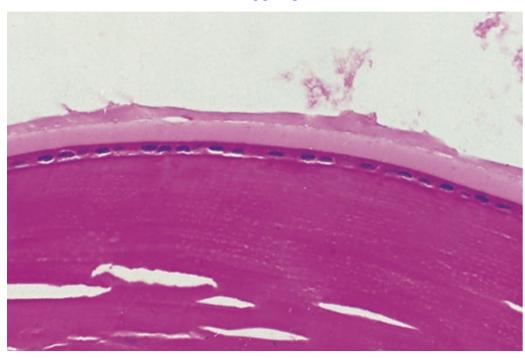
--It's deep to the 'Bowman's' (K epi is on top of Bowman's)

So it ain't cornea.

There's only one structure that has a membrane (with an epi *under* it) that surrounds amorphous material: The lens. *Is this the anterior or posterior lens?* Anterior. How can you tell?



Anterior



First: What tissue/structure is this?

It kinda looks like the anterior cornea—you can see what looks like stroma, a layer of epithelium, and Bowman's. But this interpretation is problematic in that:
--It's a single -layered epi (K epi is multi - layered)

--It's deep to the 'Bowman's' (K epi is on top of Bowman's)

So it ain't cornea.

There's only one structure that has a membrane (with an epi *under* it) that surrounds amorphous material: The lens.

Is this the anterior or posterior lens? Anterior.

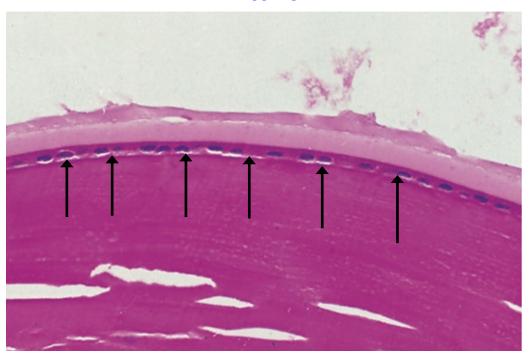
How can you tell? By the presence of

two words

(there are none posteriorly)



Anterior



First: What tissue/structure is this?

It kinda looks like the anterior cornea—you can see what looks like stroma, a layer of epithelium, and Bowman's. But this interpretation is problematic in that:
--It's a single -layered epi (K epi is multi - layered)

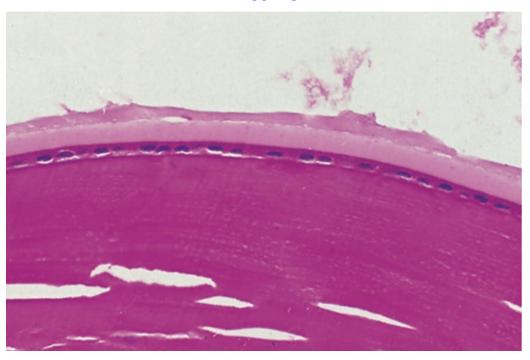
--It's deep to the 'Bowman's' (K epi is on top of Bowman's)

So it ain't cornea.

There's only one structure that has a membrane (with an epi *under* it) that surrounds amorphous material: The lens. *Is this the anterior or posterior lens?* Anterior. *How can you tell?* By the presence of epithelial cells (there are none posteriorly)



Anterior



First: What tissue/structure is this?

It kinda looks like the anterior cornea—you can see what looks like stroma, a layer of epithelium, and Bowman's. But this interpretation is problematic in that:
--It's a single -layered epi (K epi is multi - layered)

--It's deep to the 'Bowman's' (K epi is on top of Bowman's)

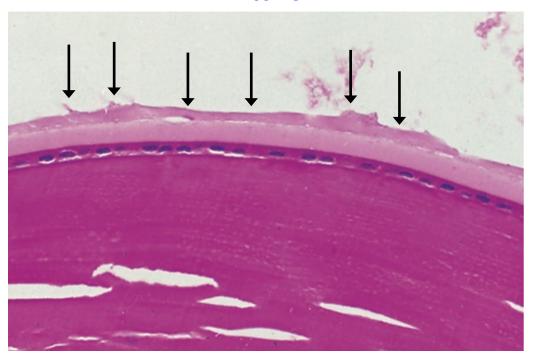
So it ain't cornea.

There's only one structure that has a membrane (with an epi *under* it) that surrounds amorphous material: The lens. *Is this the anterior or posterior lens?* Anterior. *How can you tell?* By the presence of epithelial cells (there are none posteriorly)

Now that we know what we're looking at, why are we looking at it, ie, what's abnormal?



Anterior



First: What tissue/structure is this?

It kinda looks like the anterior cornea—you can see what looks like stroma, a layer of epithelium, and Bowman's. But this interpretation is problematic in that:
--It's a single -layered epi (K epi is multi - layered)

--It's deep to the 'Bowman's' (K epi is on top of Bowman's)

So it ain't cornea.

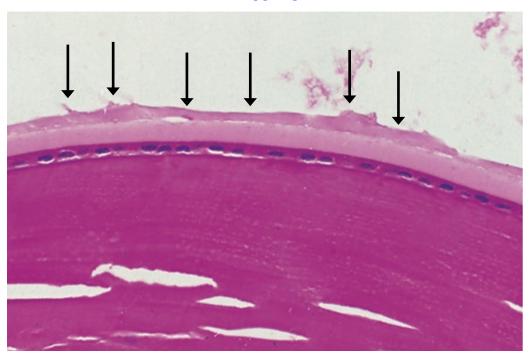
There's only one structure that has a membrane (with an epi *under* it) that surrounds amorphous material: The lens. *Is this the anterior or posterior lens?* Anterior. How can you tell? By the presence of epithelial cells (there are none posteriorly)

Now that we know what we're looking at, why are we looking at it, ie, what's abnormal?

This material on the lens proper name for



Anterior



First: What tissue/structure is this?

It kinda looks like the anterior cornea—you can see what looks like stroma, a layer of epithelium, and Bowman's. But this interpretation is problematic in that:
--It's a single -layered epi (K epi is multi - layered)

--It's deep to the 'Bowman's' (K epi is on top of Bowman's)

So it ain't cornea.

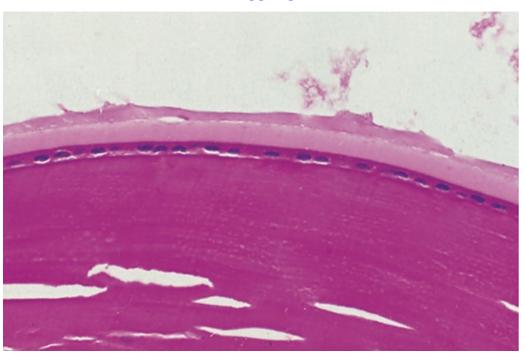
There's only one structure that has a membrane (with an epi *under* it) that surrounds amorphous material: The lens. *Is this the anterior or posterior lens?* Anterior. *How can you tell?* By the presence of epithelial cells (there are none posteriorly)

Now that we know what we're looking at, why are we looking at it, ie, what's abnormal?

This material on the lens capsule.



Anterior



What's the diagnosis?

First: What tissue/structure is this?

It kinda looks like the anterior cornea—you can see what looks like stroma, a layer of epithelium, and Bowman's. But this interpretation is problematic in that:
--It's a single -layered epi (K epi is multi - layered)

--It's deep to the 'Bowman's' (K epi is on top of Bowman's)

So it ain't cornea.

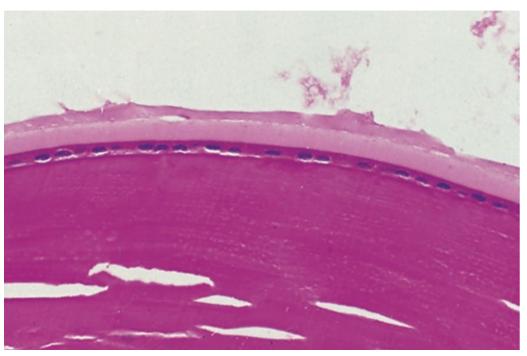
There's only one structure that has a membrane (with an epi *under* it) that surrounds amorphous material: The lens. *Is this the anterior or posterior lens?* Anterior. How can you tell? By the presence of epithelial cells (there are none posteriorly)

Now that we know what we're looking at, why are we looking at it, ie, what's abnormal?

This material on the lens capsule.



Anterior



What's the diagnosis?

First: What tissue/structure is this?
It kinda looks like the anterior cornea—you can see what looks like stroma, a layer of epithelium, and Bowman's. But this interpretation is problematic in that:
--It's a single-layered epi (K epi is multi-layered)

--It's deep to the 'Bowman's' (K epi is on top of Bowman's)

So it ain't cornea.

There's only one structure that has a membrane (with an epi *under* it) that surrounds amorphous material: The lens. *Is this the anterior or posterior lens?* Anterior. *How can you tell?* By the presence of epithelial cells (there are none posteriorly)

Now that we know what we're looking at, why are we looking at it, ie, what's abnormal?

This material on the lens capsule.

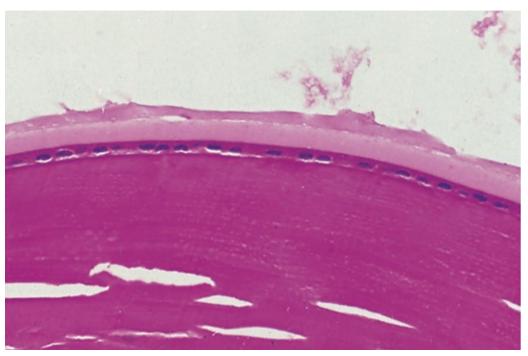
Pseudoexfoliation syndrome (PXS) is a

systemic *vs* ophthalmic

condition



Anterior



What's the diagnosis?

First: What tissue/structure is this?
It kinda looks like the anterior cornea—you can see what looks like stroma, a layer of epithelium, and Bowman's. But this interpretation is problematic in that:
--It's a single-layered epi (K epi is multi-layered)

--It's deep to the 'Bowman's' (K epi is on top of Bowman's)

So it ain't cornea.

There's only one structure that has a membrane (with an epi *under* it) that surrounds amorphous material: The lens. *Is this the anterior or posterior lens?* Anterior. *How can you tell?* By the presence of epithelial cells (there are none posteriorly)

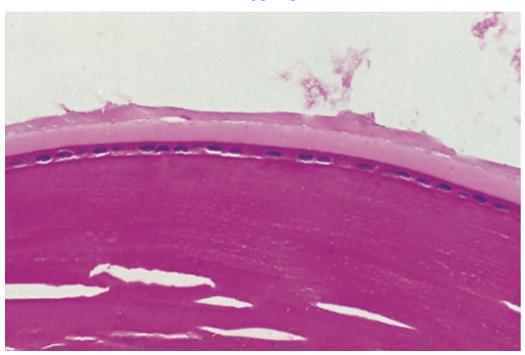
Now that we know what we're looking at, why are we looking at it, ie, what's abnormal?

This material on the lens capsule.

Pseudoexfoliation syndrome (PXS) is a systemic condition



Anterior



What's the diagnosis?

First: What tissue/structure is this?
It kinda looks like the anterior cornea—you can see what looks like stroma, a layer of epithelium, and Bowman's. But this interpretation is problematic in that:
--It's a single-layered epi (K epi is multi-layered)

--It's deep to the 'Bowman's' (K epi is on top of Bowman's)

So it ain't cornea.

There's only one structure that has a membrane (with an epi *under* it) that surrounds amorphous material: The lens. *Is this the anterior or posterior lens?* Anterior. *How can you tell?* By the presence of epithelial cells (there are none posteriorly)

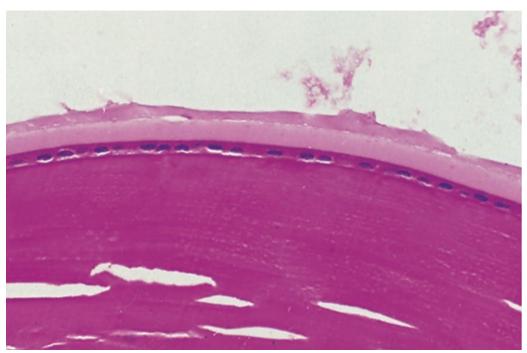
Now that we know what we're looking at, why are we looking at it, ie, what's abnormal?

This material on the lens capsule.

Pseudoexfoliation syndrome (PXS) is a systemic condition characterized by the production and accumulation of material throughout the anterior segment*, including upon the lens capsule.



Anterior



What's the diagnosis?

First: What tissue/structure is this?
It kinda looks like the anterior cornea—you can see what looks like stroma, a layer of epithelium, and Bowman's. But this interpretation is problematic in that:
--It's a single-layered epi (K epi is multi-layered)

--It's deep to the 'Bowman's' (K epi is on top of Bowman's)

So it ain't cornea.

There's only one structure that has a membrane (with an epi *under* it) that surrounds amorphous material: The lens. *Is this the anterior or posterior lens?* Anterior. *How can you tell?* By the presence of epithelial cells (there are none posteriorly)

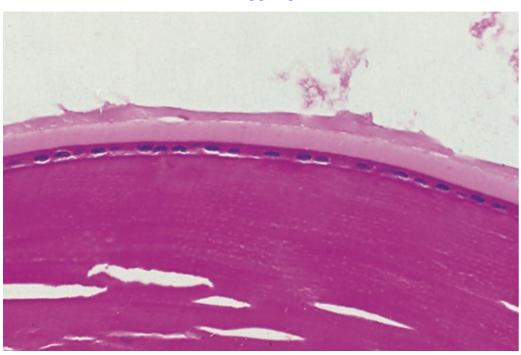
Now that we know what we're looking at, why are we looking at it, ie, what's abnormal?

This material on the lens capsule.

Pseudoexfoliation syndrome (PXS) is a systemic condition characterized by the production and accumulation of fibrillar material throughout the anterior segment*, including upon the lens capsule.



Anterior



What's the diagnosis?

First: What tissue/structure is this?
It kinda looks like the anterior cornea—you can see what looks like stroma, a layer of epithelium, and Bowman's. But this interpretation is problematic in that:
--It's a single-layered epi (K epi is multi-layered)

--It's deep to the 'Bowman's' (K epi is on top of Bowman's)

So it ain't cornea.

There's only one structure that has a membrane (with an epi *under* it) that surrounds amorphous material: The lens. *Is this the anterior or posterior lens?* Anterior. *How can you tell?* By the presence of epithelial cells (there are none posteriorly)

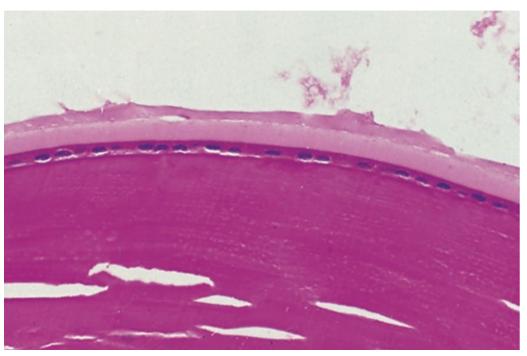
Now that we know what we're looking at, why are we looking at it, ie, what's abnormal?

This material on the lens capsule.

Pseudoexfoliation syndrome (PXS) is a systemic condition characterized by the production and accumulation of fibrillar material throughout the anterior segment*, including upon the lens capsule. It is associated with ophthalmic condition



Anterior



What's the diagnosis?

First: What tissue/structure is this?
It kinda looks like the anterior cornea—you can see what looks like stroma, a layer of epithelium, and Bowman's. But this interpretation is problematic in that:
--It's a single-layered epi (K epi is multi-layered)

--It's deep to the 'Bowman's' (K epi is on top of Bowman's)

So it ain't cornea.

There's only one structure that has a membrane (with an epi *under* it) that surrounds amorphous material: The lens. *Is this the anterior or posterior lens?* Anterior. *How can you tell?* By the presence of epithelial cells (there are none posteriorly)

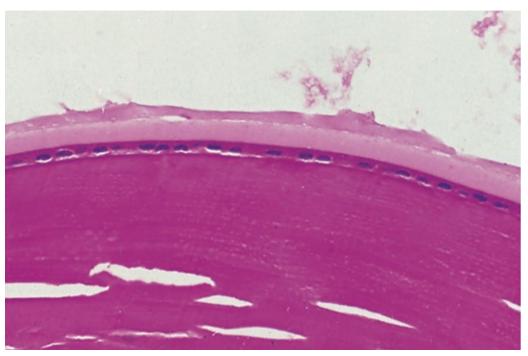
Now that we know what we're looking at, why are we looking at it, ie, what's abnormal?

This material on the lens capsule.

Pseudoexfoliation syndrome (PXS) is a systemic condition characterized by the production and accumulation of fibrillar material throughout the anterior segment*, including upon the lens capsule. It is associated with glaucoma*



Anterior



What's the diagnosis?

First: What tissue/structure is this?
It kinda looks like the anterior cornea—you can see what looks like stroma, a layer of epithelium, and Bowman's. But this interpretation is problematic in that:
--It's a single-layered epi (K epi is multi-layered)

--It's deep to the 'Bowman's' (K epi is on top of Bowman's)

So it ain't cornea.

There's only one structure that has a membrane (with an epi *under* it) that surrounds amorphous material: The lens. *Is this the anterior or posterior lens?* Anterior. *How can you tell?* By the presence of epithelial cells (there are none posteriorly)

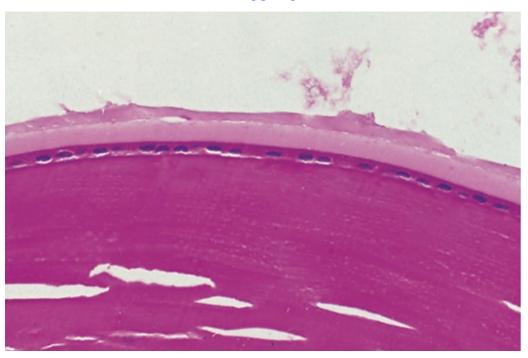
Now that we know what we're looking at, why are we looking at it, ie, what's abnormal?

This material on the lens capsule.

Pseudoexfoliation syndrome (PXS) is a systemic condition characterized by the production and accumulation of fibrillar material throughout the anterior segment*, including upon the lens capsule. It is associated with glaucoma*



Anterior



What's the diagnosis?

First: What tissue/structure is this?
It kinda looks like the anterior cornea—you can see what looks like stroma, a layer of epithelium, and Bowman's. But this interpretation is problematic in that:
--It's a single-layered epi (K epi is multi-layered)

--It's deep to the 'Bowman's' (K epi is on top of Bowman's)

So it ain't cornea.

There's only one structure that has a membrane (with an epi *under* it) that surrounds amorphous material: The lens. *Is this the anterior or posterior lens?* Anterior. *How can you tell?* By the presence of epithelial cells (there are none posteriorly)

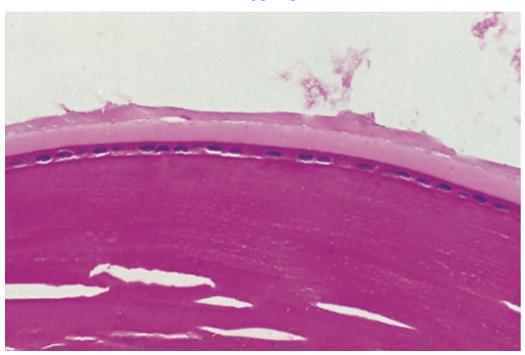
Now that we know what we're looking at, why are we looking at it, ie, what's abnormal?

This material on the lens capsule.

Pseudoexfoliation syndrome (PXS) is a systemic condition characterized by the production and accumulation of fibrillar material throughout the anterior segment*, including upon the lens capsule. It is associated with glaucoma*



Anterior



What's the diagnosis?

First: What tissue/structure is this?
It kinda looks like the anterior cornea—you can see what looks like stroma, a layer of epithelium, and Bowman's. But this interpretation is problematic in that:
--It's a single-layered epi (K epi is multi-layered)

--It's deep to the 'Bowman's' (K epi is on top of Bowman's)

So it ain't cornea.

There's only one structure that has a membrane (with an epi *under* it) that surrounds amorphous material: The lens. *Is this the anterior or posterior lens?* Anterior. *How can you tell?* By the presence of epithelial cells (there are none posteriorly)

Now that we know what we're looking at, why are we looking at it, ie, what's abnormal?

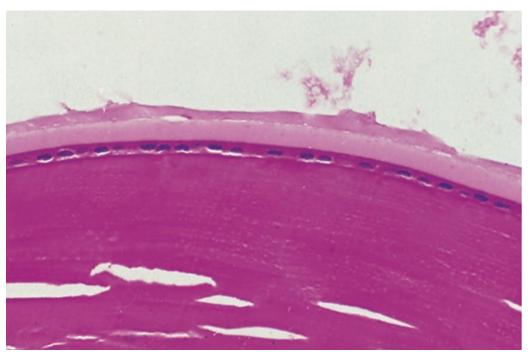
This material on the lens capsule.

Pseudoexfoliation syndrome (PXS) is a systemic condition characterized by the production and accumulation of fibrillar material throughout the anterior segment*, including upon the lens capsule. It is associated with glaucoma*

^{*}Specifically *open* -angle glaucoma, specifically *secondary* open-angle glaucoma.



Anterior



What's the diagnosis?

First: What tissue/structure is this?
It kinda looks like the anterior cornea—you can see what looks like stroma, a layer of epithelium, and Bowman's. But this interpretation is problematic in that:
--It's a single-layered epi (K epi is multi-layered)

--It's deep to the 'Bowman's' (K epi is on top of Bowman's)

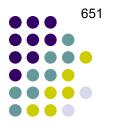
So it ain't cornea.

There's only one structure that has a membrane (with an epi *under* it) that surrounds amorphous material: The lens. *Is this the anterior or posterior lens?* Anterior. *How can you tell?* By the presence of epithelial cells (there are none posteriorly)

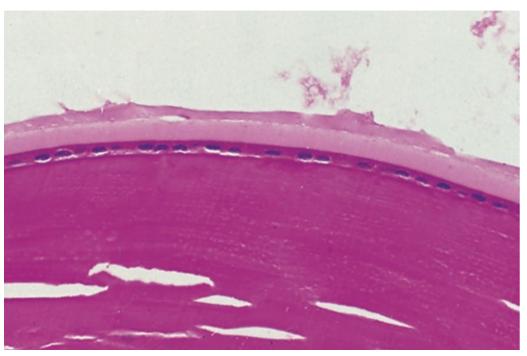
Now that we know what we're looking at, why are we looking at it, ie, what's abnormal?

This material on the lens capsule.

Pseudoexfoliation syndrome (PXS) is a systemic condition characterized by the production and accumulation of fibrillar material throughout the anterior segment*, including upon the lens capsule. It is associated with glaucoma* and an increased risk of intra-op complications during surgery.



Anterior



What's the diagnosis?

First: What tissue/structure is this?
It kinda looks like the anterior cornea—you can see what looks like stroma, a layer of epithelium, and Bowman's. But this interpretation is problematic in that:
--It's a single-layered epi (K epi is multi-layered)

--It's deep to the 'Bowman's' (K epi is on top of Bowman's)

So it ain't cornea.

There's only one structure that has a membrane (with an epi *under* it) that surrounds amorphous material: The lens. *Is this the anterior or posterior lens?* Anterior. *How can you tell?* By the presence of epithelial cells (there are none posteriorly)

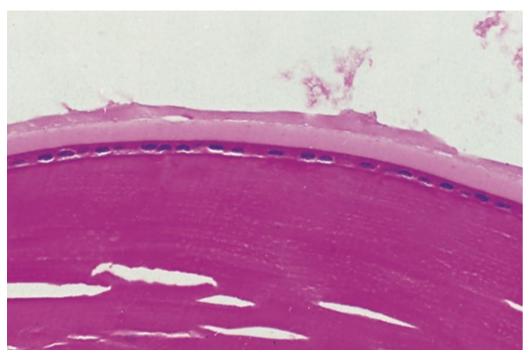
Now that we know what we're looking at, why are we looking at it, ie, what's abnormal?

This material on the lens capsule.

Pseudoexfoliation syndrome (PXS) is a systemic condition characterized by the production and accumulation of fibrillar material throughout the anterior segment*, including upon the lens capsule. It is associated with glaucoma* and an increased risk of intra-op complications during cataract surgery.



Anterior



What's the diagnosis?

First: What tissue/structure is this?
It kinda looks like the anterior cornea—you can see what looks like stroma, a layer of epithelium, and Bowman's. But this interpretation is problematic in that:
--It's a single-layered epi (K epi is multi-layered)

--It's deep to the 'Bowman's' (K epi is on top of Bowman's)

So it ain't cornea.

There's only one structure that has a membrane (with an epi *under* it) that surrounds amorphous material: The lens. *Is this the anterior or posterior lens?* Anterior. *How can you tell?* By the presence of epithelial cells (there are none posteriorly)

Now that we know what we're looking at, why are we looking at it, ie, what's abnormal?

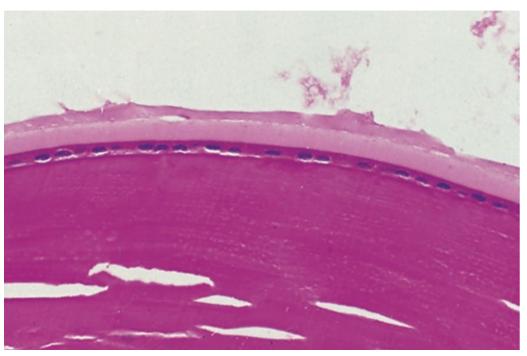
This material on the lens capsule.

Pseudoexfoliation syndrome (PXS) is a systemic condition characterized by the production and accumulation of fibrillar material throughout the anterior segment*, including upon the lens capsule. It is associated with glaucoma* and an increased risk of intra-op complications during cataract surgery.

The appearance of the accumulated material has been likened to a buzzword pattern and 'buzz-term (humands)'.



Anterior



What's the diagnosis?

First: What tissue/structure is this?
It kinda looks like the anterior cornea—you can see what looks like stroma, a layer of epithelium, and Bowman's. But this interpretation is problematic in that:
--It's a single-layered epi (K epi is multi-layered)

--It's deep to the 'Bowman's' (K epi is on top of Bowman's)

So it ain't cornea.

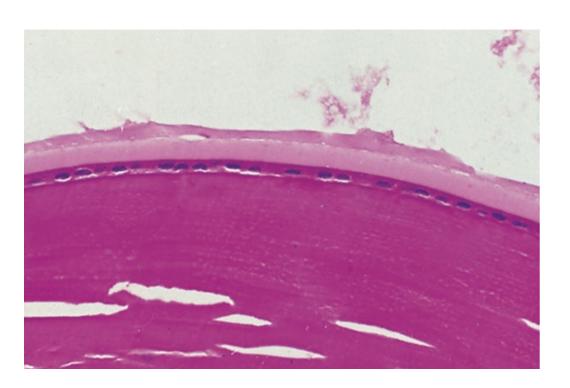
There's only one structure that has a membrane (with an epi *under* it) that surrounds amorphous material: The lens. *Is this the anterior or posterior lens?* Anterior. *How can you tell?* By the presence of epithelial cells (there are none posteriorly)

Now that we know what we're looking at, why are we looking at it, ie, what's abnormal?

This material on the lens capsule.

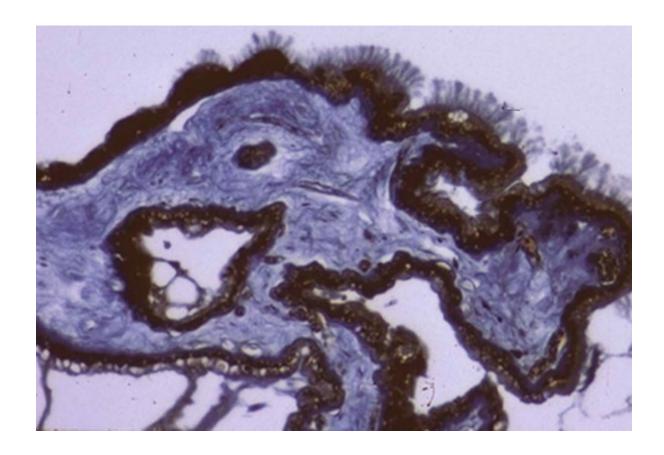
Pseudoexfoliation syndrome (PXS) is a systemic condition characterized by the production and accumulation of fibrillar material throughout the anterior segment*, including upon the lens capsule. It is associated with glaucoma* and an increased risk of intra-op complications during cataract surgery. The appearance of the accumulated material has been likened to a sawtooth pattern and ' iron filings '.





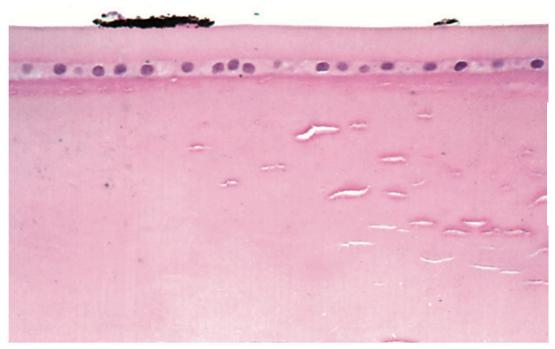
Keeping it 100: This is a *terrible* pic to use when asserting that PXS fibrillar material adopts what could be called an 'iron filings' configuration.





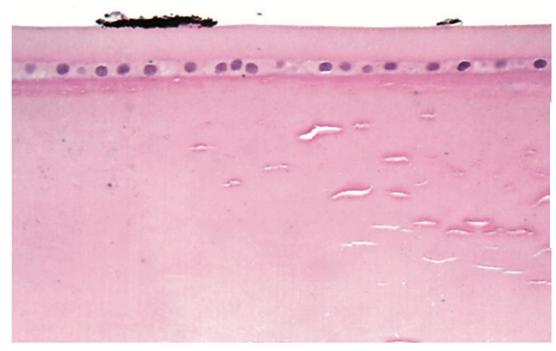
Keeping it 100: This is a *terrible* pic to use when asserting that PXS fibrillar material adopts what could be called an 'iron filings' configuration. **This** pic of fibrillar material on the ciliary body does justice to the assertion. Just FYI.





Once again, we're faced with (what we now recognize as) an image of the anterior aspect of the lens. And once again, the lens capsule has some gradeau on it, only instead of pinkish and sawtooth-y it's black and flat.

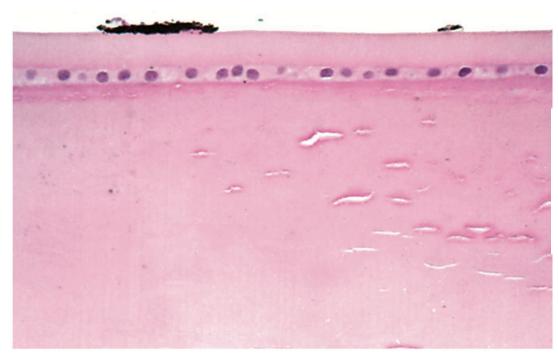




Once again, we're faced with (what we now recognize as) an image of the anterior aspect of the lens. And once again, the lens capsule has some gradeau on it, only instead of pinkish and sawtooth-y it's black and flat. What's going on here?

What's the diagnosis?





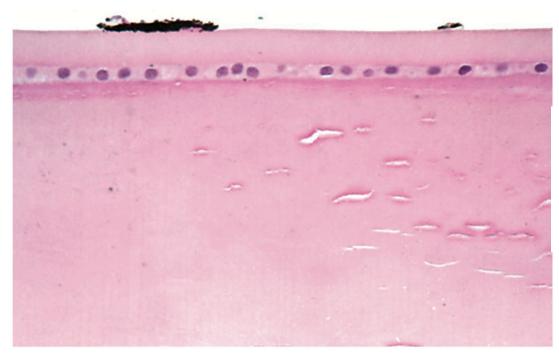
Once again, we're faced with (what we now recognize as) an image of the anterior aspect of the lens. And once again, the lens capsule has some gradeau on it, only instead of pinkish and sawtooth-y it's black and flat. What's going on here?

What's the diagnosis?

Pigment dispersion syndrome (PDS) results from excessive contact between the surface and the

iris



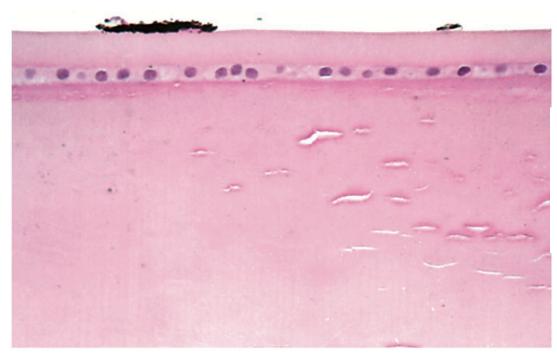


Once again, we're faced with (what we now recognize as) an image of the anterior aspect of the lens. And once again, the lens capsule has some gradeau on it, only instead of pinkish and sawtooth-y it's black and flat. What's going on here?

What's the diagnosis?

Pigment dispersion syndrome (PDS) results from excessive contact between the posterior iris surface and the zonules.



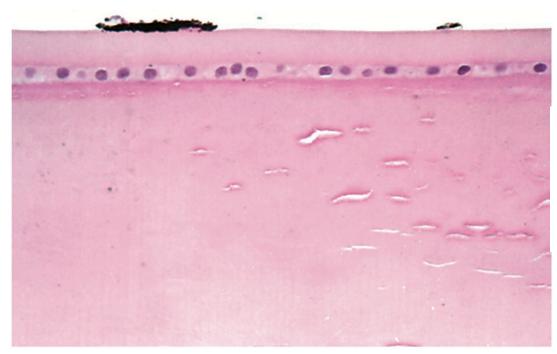


Once again, we're faced with (what we now recognize as) an image of the anterior aspect of the lens. And once again, the lens capsule has some gradeau on it, only instead of pinkish and sawtooth-y it's black and flat. What's going on here?

What's the diagnosis?

Pigment dispersion syndrome (PDS) results from excessive contact between the posterior iris surface and the zonules. Friction between these structures liberates two words from the iris



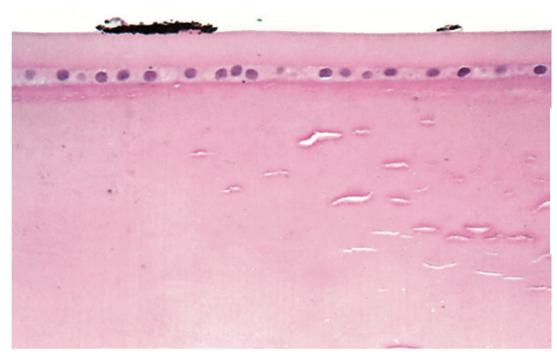


Once again, we're faced with (what we now recognize as) an image of the anterior aspect of the lens. And once again, the lens capsule has some gradeau on it, only instead of pinkish and sawtooth-y it's black and flat. What's going on here?

What's the diagnosis?

Pigment dispersion syndrome (PDS) results from excessive contact between the posterior iris surface and the zonules. Friction between these structures liberates pigment granules from the iris



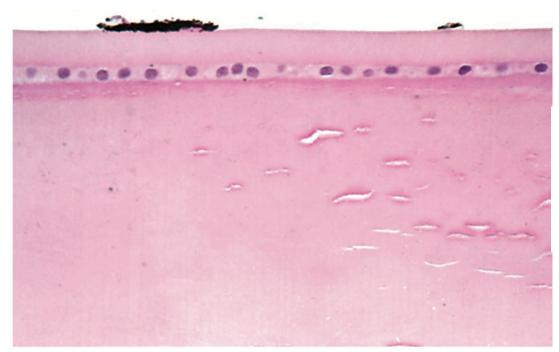


Once again, we're faced with (what we now recognize as) an image of the anterior aspect of the lens. And once again, the lens capsule has some gradeau on it, only instead of pinkish and sawtooth-y it's black and flat. What's going on here?

What's the diagnosis?

Pigment dispersion syndrome (PDS) results from excessive contact between the posterior iris surface and the zonules. Friction between these structures liberates pigment granules from the iris, which subsequently deposit on intraocular structures including the lens capsule.



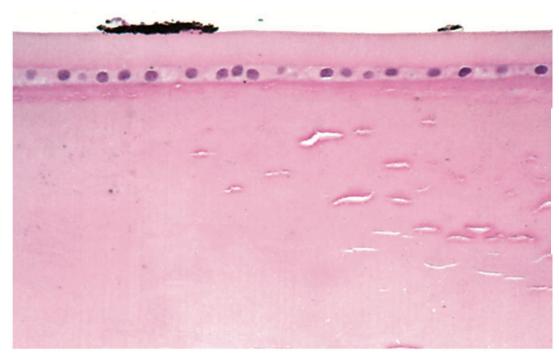


Once again, we're faced with (what we now recognize as) an image of the anterior aspect of the lens. And once again, the lens capsule has some gradeau on it, only instead of pinkish and sawtooth-y it's black and flat. What's going on here?

What's the diagnosis?

Pigment dispersion syndrome (PDS) results from excessive contact between the posterior iris surface and the zonules. Friction between these structures liberates pigment granules from the iris, which subsequently deposit on intraocular structures including the lens capsule. Pts with PDS are at risk of developing

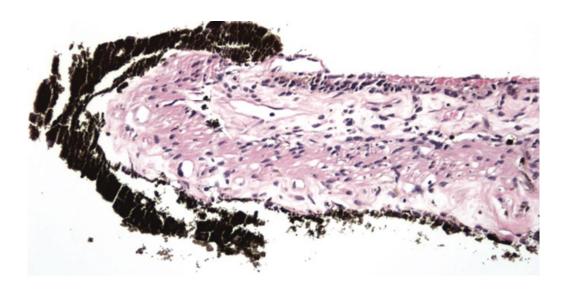




Once again, we're faced with (what we now recognize as) an image of the anterior aspect of the lens. And once again, the lens capsule has some gradeax on it, only instead of pinkish and sawtooth-y it's black and flat. What's going on here?

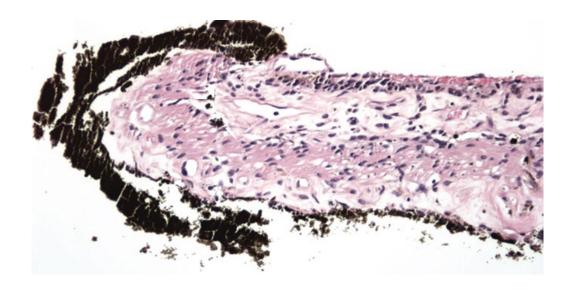
What's the diagnosis?

Pigment dispersion syndrome (PDS) results from excessive contact between the posterior iris surface and the zonules. Friction between these structures liberates pigment granules from the iris, which subsequently deposit on intraocular structures including the lens capsule. Pts with PDS are at risk of developing pigmentary glaucoma.



First things first: What tissue is this?

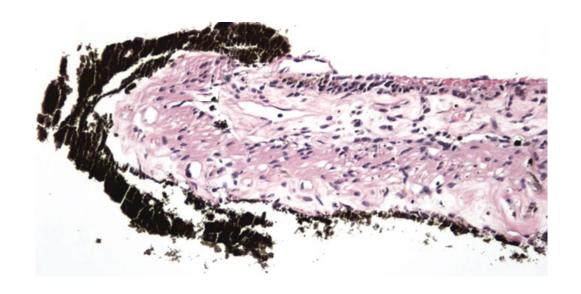




First things first: What tissue is this? It's long and skinny, and has a natural end—is it a section of eyelid at the margin?



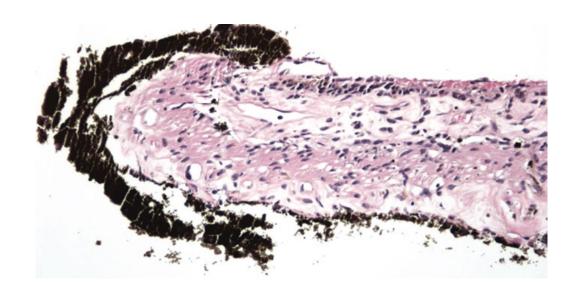




First things first: What tissue is this?

It's long and skinny, and has a natural end—is it a section of eyelid at the margin? No, because it lacks any semblance of skin/conj structures, and the end looks **nothing** like a lid margin.

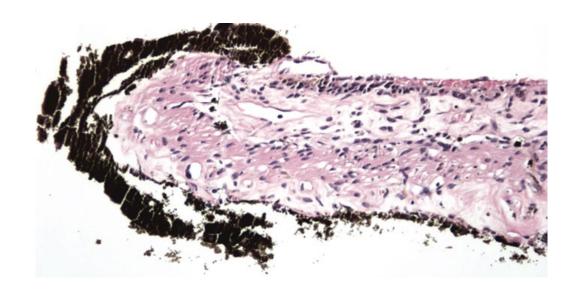




First things first: What tissue is this?

It's long and skinny, and has a natural end—is it a section of eyelid at the margin? No, because it lacks any semblance of skin/conj structures, and the end looks **nothing** like a lid margin. There's only one non-lid structure shaped like this—what is it?

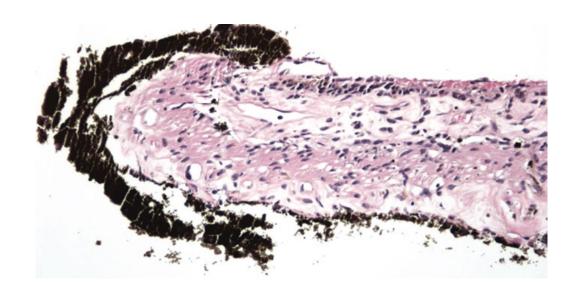




First things first: What tissue is this?

It's long and skinny, and has a natural end—is it a section of eyelid at the margin? No, because it lacks any semblance of skin/conj structures, and the end looks **nothing** like a lid margin. There's only one non-lid structure shaped like this—what is it? The iris.





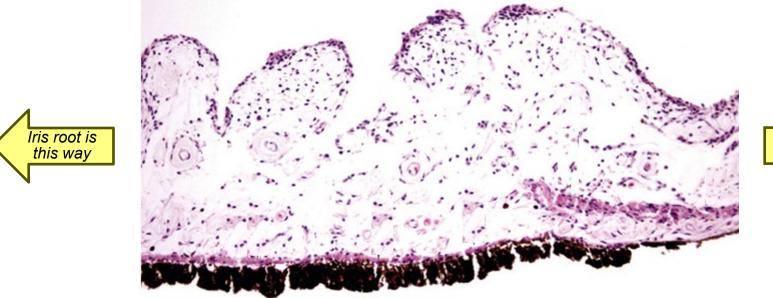
First things first: What tissue is this?

It's long and skinny, and has a natural end—is it a section of eyelid at the margin? No, because it lacks any semblance of skin/conj structures, and the end looks **nothing** like a lid margin. There's only one non-lid structure shaped like this—what is it? The iris.

Before getting into the pathology here, let's review normal iris anatomy...



Anterior? Posterior?



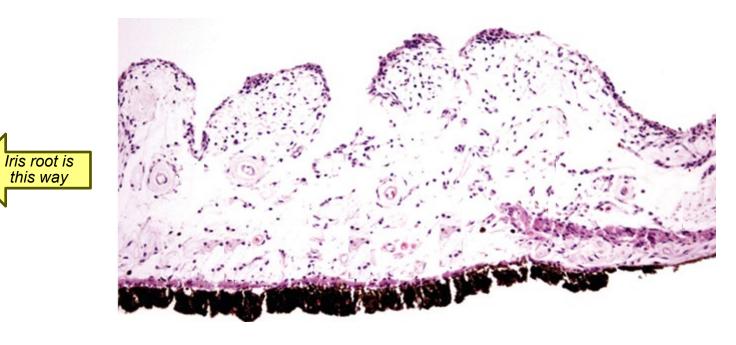


Anterior? Posterior?

Normal iris. Which surface is anterior, and which is posterior?



Anterior

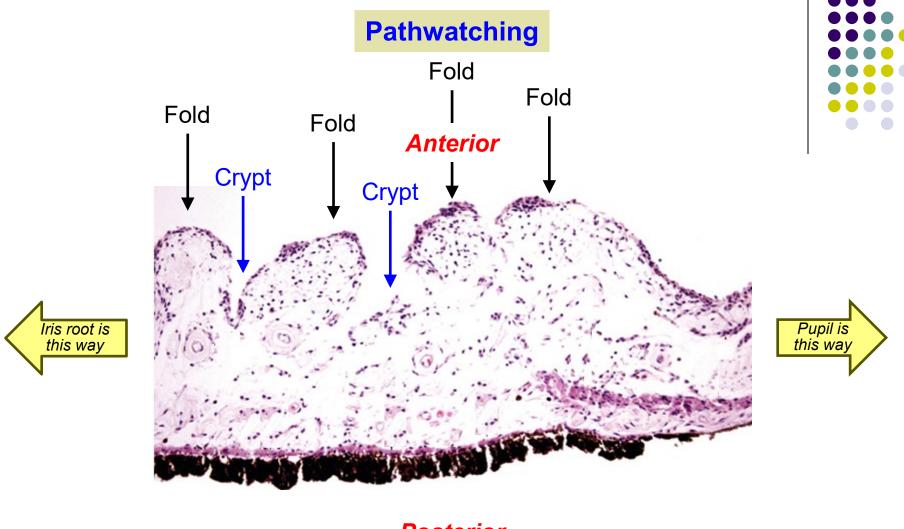


Pupil is this way

Posterior

Normal iris. Which surface is anterior, and which is posterior?

The anterior surface is corrugated, with large separated by deep



673

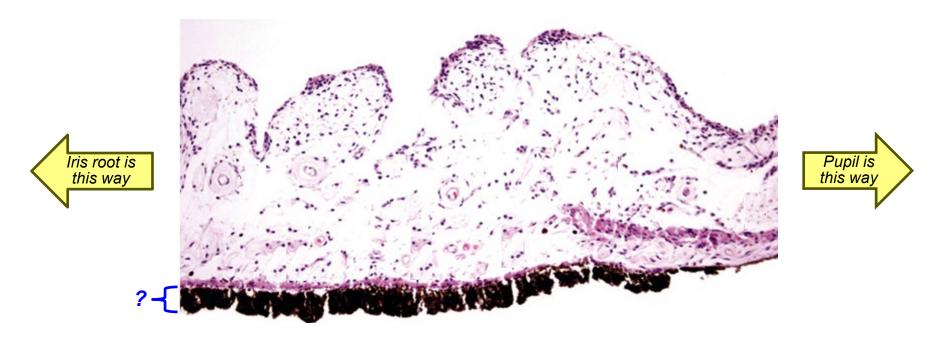
Posterior

Normal iris. Which surface is anterior, and which is posterior?

The anterior surface is corrugated, with large folds separated by deep crypts.



Anterior



Posterior

Normal iris. Which surface is anterior, and which is posterior?

The anterior surface is corrugated, with large folds separated by deep crypts. The posterior surface

is lined by a densely-pigmented

layer of

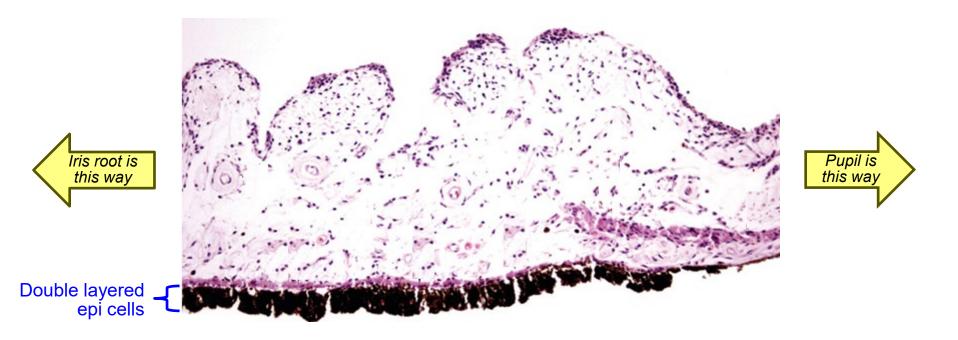
cell type

cells oriented

base-to-base base-to-apex?

675

Anterior

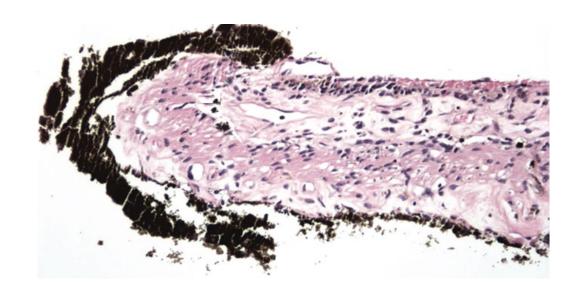


Posterior

Normal iris. Which surface is anterior, and which is posterior?

The anterior surface is corrugated, with large folds separated by deep crypts. The posterior surface is lined by a densely-pigmented double layer of epithelial cells oriented apex-to-apex.*





First things first: What tissue is this?

It's long and skinny, and has a natural end—is it a section of eyelid at the margin? No, because it lacks any semblance of skin/conj structures, and the end looks **nothing** like a lid margin. There's only one non-lid structure shaped like this—what is it? The iris.

Before getting into the pathology here, let's review normal iris anatomy...

Now back to our path slide.

Pathwatching Anterior To iris root Pupil Posterior

677

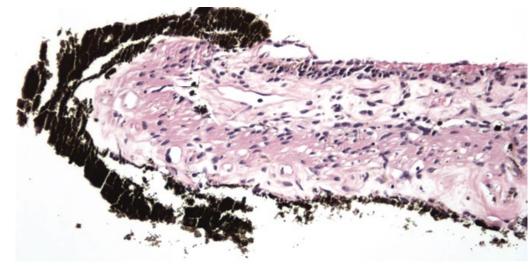
First things first: What tissue is this?

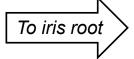
It's long and skinny, and has a natural end—is it a section of eyelid at the margin? No, because it lacks any semblance of skin/conj structures, and the end looks **nothing** like a lid margin. There's only one non-lid structure shaped like this—what is it? The iris.

Before getting into the pathology here, let's review normal iris anatomy...

Now back to our path slide. First, here is the orientation of what we're looking at.

Anterior





678

Posterior

First things first: What tissue is this?

Pupil

It's long and skinny, and has a natural end—is it a section of eyelid at the margin? No, because it lacks any semblance of skin/conj structures, and the end looks **nothing** like a lid margin. There's only one non-lid structure shaped like this—what is it? The iris.

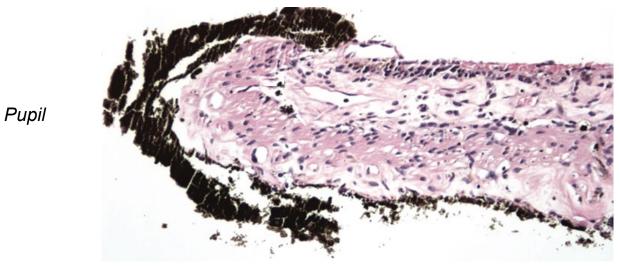
Before getting into the pathology here, let's review normal iris anatomy...

Now back to our path slide. First, here is the orientation of what we're looking at. Now, based on our new knowledge of normal iris anatomy we can see a couple of problems right off the bat:

--?

--?

Anterior





679

Posterior

First things first: What tissue is this?

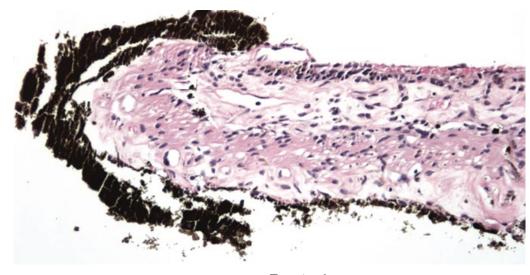
It's long and skinny, and has a natural end—is it a section of eyelid at the margin? No, because it lacks any semblance of skin/conj structures, and the end looks **nothing** like a lid margin. There's only one non-lid structure shaped like this—what is it? The iris.

Before getting into the pathology here, let's review normal iris anatomy...

Now back to our path slide. First, here is the orientation of what we're looking at. Now, based on our new knowledge of normal iris anatomy we can see a couple of problems right off the bat:

--The anterior iris is unnaturally

Anterior





680

Posterior

First things first: What tissue is this?

Pupil

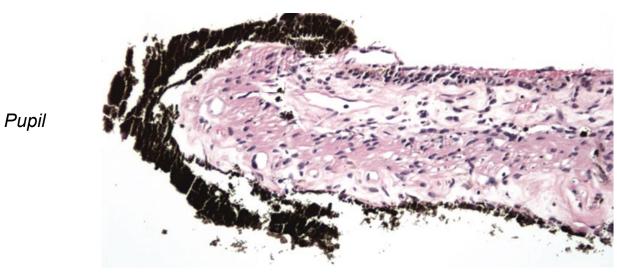
It's long and skinny, and has a natural end—is it a section of eyelid at the margin? No, because it lacks any semblance of skin/conj structures, and the end looks **nothing** like a lid margin. There's only one non-lid structure shaped like this—what is it? The iris.

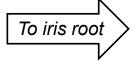
Before getting into the pathology here, let's review normal iris anatomy...

Now back to our path slide. First, here is the orientation of what we're looking at. Now, based on our new knowledge of normal iris anatomy we can see a couple of problems right off the bat:

--The anterior iris is unnaturally flat

Anterior





681

Posterior

First things first: What tissue is this?

It's long and skinny, and has a natural end—is it a section of eyelid at the margin? No, because it lacks any semblance of skin/conj structures, and the end looks **nothing** like a lid margin. There's only one non-lid structure shaped like this—what is it? The iris.

Before getting into the pathology here, let's review normal iris anatomy...

Now back to our path slide. First, here is the orientation of what we're looking at. Now, based on our new knowledge of normal iris anatomy we can see a couple of problems right off the bat:

- -- The anterior iris is unnaturally flat
- --The posterior pigmented epi bilayer has

lotsa words

Pathwatching Anterior To iris root Pupil

682

First things first: What tissue is this?

It's long and skinny, and has a natural end—is it a section of eyelid at the margin? No, because it lacks any semblance of skin/conj structures, and the end looks **nothing** like a lid margin. There's only one non-lid structure shaped like this—what is it? The iris.

Posterior

Before getting into the pathology here, let's review normal iris anatomy...

Now back to our path slide. First, here is the orientation of what we're looking at. Now, based on our new knowledge of normal iris anatomy we can see a couple of problems right off the bat:

- -- The anterior iris is unnaturally flat
- --The posterior pigmented epi bilayer has come around the pupil margin onto the anterior iris.

683 **Pathwatching** Anterior To iris root Pupil

What's the diagnosis?

First things first: What tissue is this?

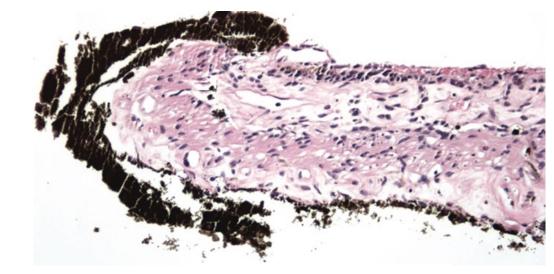
It's long and skinny, and has a natural end—is it a section of eyelid at the margin? No, because it lacks any semblance of skin/conj structures, and the end looks **nothing** like a lid margin. There's only one non-lid structure shaped like this—what is it? The iris.

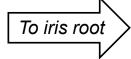
Before getting into the pathology here, let's review normal iris anatomy...

Now back to our path slide. First, here is the orientation of what we're looking at. Now, based on our new knowledge of normal iris anatomy we can see a couple of problems right off the bat:

- -- The anterior iris is unnaturally flat
- --The posterior pigmented epi bilayer has come around the pupil margin onto the anterior iris. When you see an iris with a flat anterior surface + pigmented epithelium coming around like this, one diagnosis should come to mind:

Anterior





What's the diagnosis?

First things Rubeosis iridis (iris neovascularization) is associated with a number of conditions.

It's long an semblance

lacks any non-lid 684

structure shaped like this—what is it? The iris.

Before getting into the pathology here, let's review normal iris anatomy...

Now back to our path slide. Based on our new knowledge of normal iris anatomy we can see a couple of problems right off the bat:

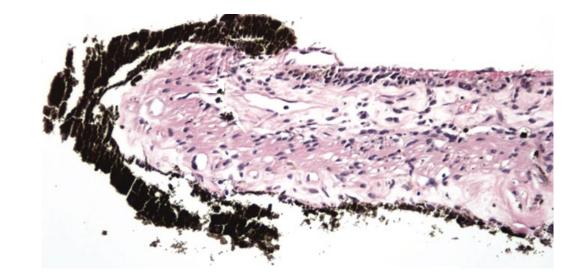
-- The anterior iris is unnaturally flat

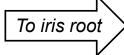
Pupil

--The posterior pigmented epi bilayer has come around the pupil margin onto the anterior iris.

When you see an iris with a flat anterior surface + pigmented epithelium coming around like this, one diagnosis should come to mind:

Anterior





What's the diagnosis?

semblance

Pupil

First things Rubeosis iridis (iris neovascularization) is associated with a number of conditions. It's long an The final pathologic pathway involves the exuberant (over)production of

lacks any non-lid

685

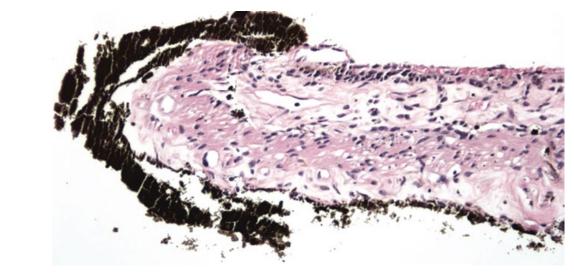
structure shaped like this—what is it? The iris.

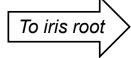
Before getting into the pathology here, let's review normal iris anatomy...

Now back to our path slide. Based on our new knowledge of normal iris anatomy we can see a couple of problems right off the bat:

- -- The anterior iris is unnaturally flat
- --The posterior pigmented epi bilayer has come around the pupil margin onto the anterior iris.

Anterior





What's the diagnosis?

semblance

Pupil

First things Rubeosis iridis (iris neovascularization) is associated with a number of conditions. It's long an The final pathologic pathway involves the exuberant (over)production of VEGF

lacks any non-lid

686

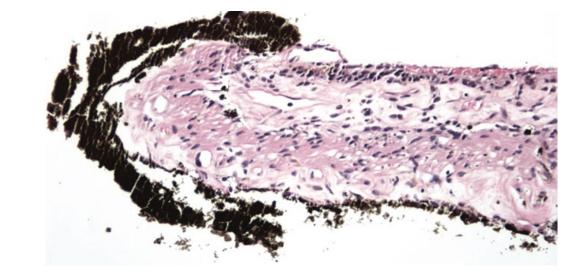
structure shaped like this—what is it? The iris.

Before getting into the pathology here, let's review normal iris anatomy...

Now back to our path slide. Based on our new knowledge of normal iris anatomy we can see a couple of problems right off the bat:

- -- The anterior iris is unnaturally flat
- --The posterior pigmented epi bilayer has come around the pupil margin onto the anterior iris.

Anterior





What's the diagnosis?

Pupil

First things Rubeosis iridis (iris neovascularization) is associated with a number of conditions. It's long an The final pathologic pathway involves the exuberant (over)production of VEGF, semblance a signaling molecule that

lacks any non-lid

687

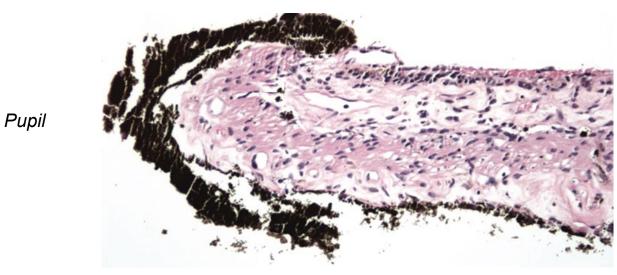
structure shaped like this—what is it? The iris.

Before getting into the pathology here, let's review normal iris anatomy...

Now back to our path slide. Based on our new knowledge of normal iris anatomy we can see a couple of problems right off the bat:

- -- The anterior iris is unnaturally flat
- --The posterior pigmented epi bilayer has come around the pupil margin onto the anterior iris.

Anterior





What's the diagnosis?

First things Rubeosis iridis (iris neovascularization) is associated with a number of conditions. It's long an The final pathologic pathway involves the exuberant (over)production of VEGF, semblance a signaling molecule that promotes neovascularization.

lacks any non-lid

688

structure shaped like this—what is it? The iris.

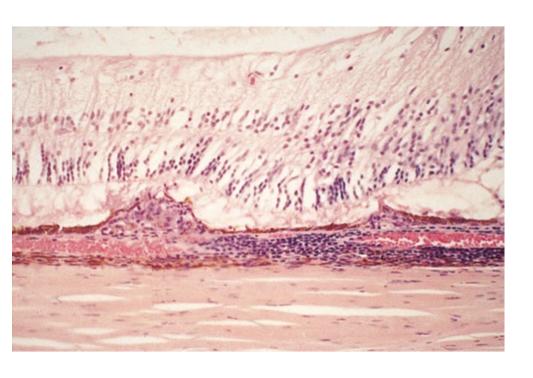
Before getting into the pathology here, let's review normal iris anatomy...

Now back to our path slide. Based on our new knowledge of normal iris anatomy we can see a couple of problems right off the bat:

- -- The anterior iris is unnaturally flat
- --The posterior pigmented epi bilayer has come around the pupil margin onto the anterior iris.

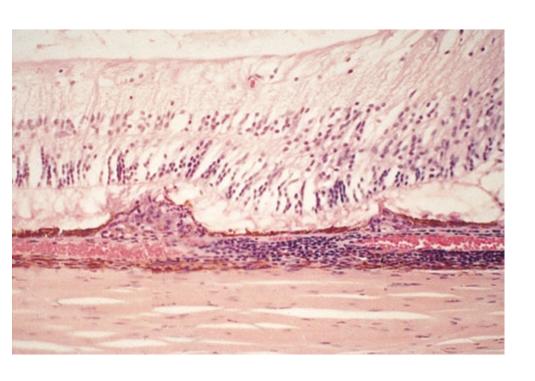


Hopefully you recognize what we're looking at here—it's the structure along with the structure and underlying last structure

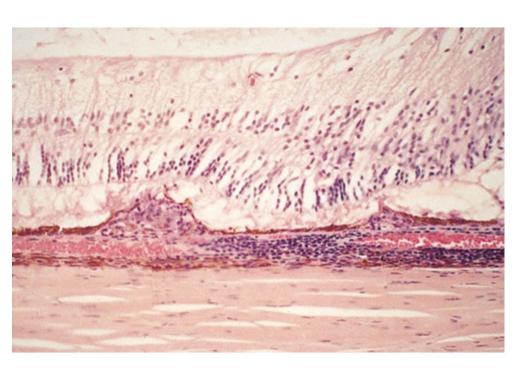




Hopefully you recognize what we're looking at here—it's the retina along with the choroid and underlying sclera.

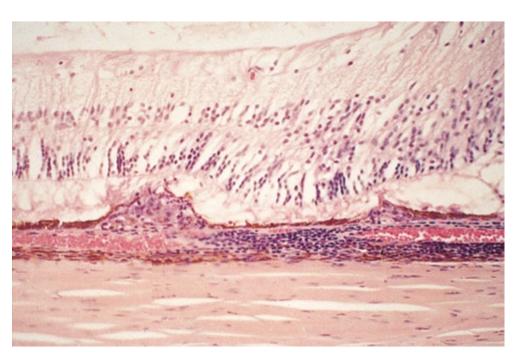






Hopefully you recognize what we're looking at here—it's the retina along with the choroid and underlying sclera.



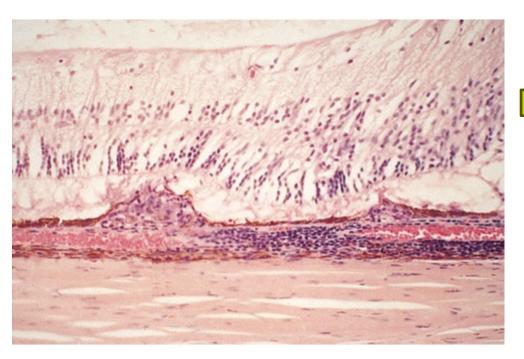


Hopefully you recognize what we're looking at here—it's the retina along with the choroid and underlying sclera.

What's going on here—what are we supposed to take note of?

--First, while I realize the neurosensory retina looks wonky, that's not the point of the slide.





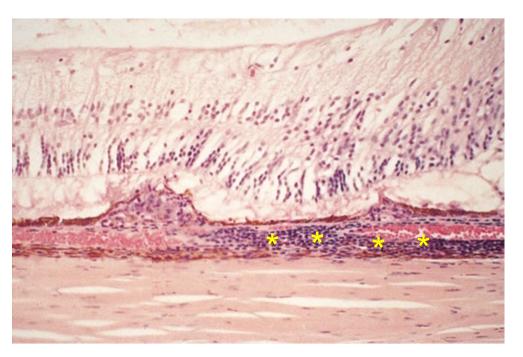
Hopefully you recognize what we're looking at here—it's the retina along with the choroid and underlying sclera.

What's going on here—what are we supposed to take note of?

--First, while I realize the neurosensory retina looks wonky, that's not the point of the slide.

--The choroid is filled with innumerable cells.

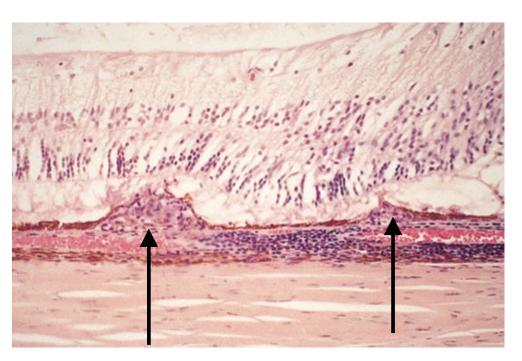




Hopefully you recognize what we're looking at here—it's the retina along with the choroid and underlying sclera.

- --First, while I realize the neurosensory retina looks wonky, that's not the point of the slide.
- --The choroid is filled with innumerable inflammatory cells. (asterisks)

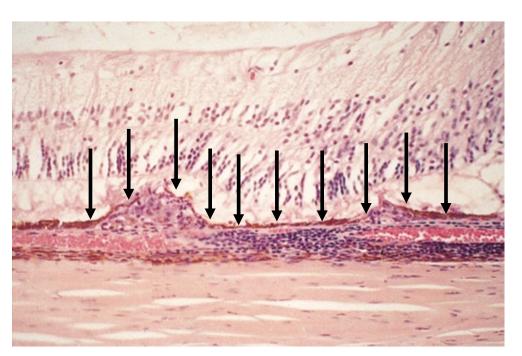




Hopefully you recognize what we're looking at here—it's the retina along with the choroid and underlying sclera.

- --First, while I realize the neurosensory retina looks wonky, that's not the point of the slide.
- --The choroid is filled with innumerable inflammatory cells.
- --Two focal aggregates of inflammatory cells can be seen **here** and **here**. Drill down on these aggregates—exactly where are they located?

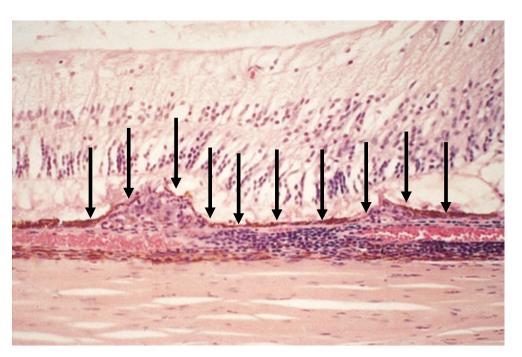




Hopefully you recognize what we're looking at here—it's the retina along with the choroid and underlying sclera.

- --First, while I realize the neurosensory retina looks wonky, that's not the point of the slide.
- --The choroid is filled with innumerable inflammatory cells.
- --Two focal aggregates of inflammatory cells can be seen **here** and **here**. Drill down on these aggregates—exactly where are they located? Note what's running across their tops—a thin, densely pigmented line. This is the

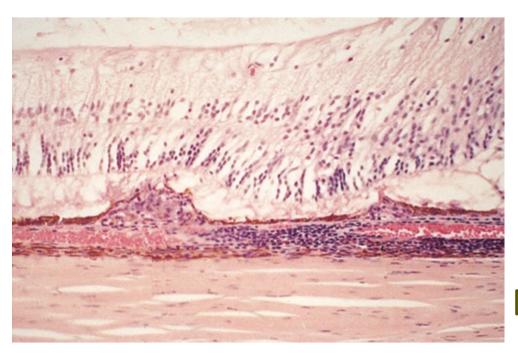




Hopefully you recognize what we're looking at here—it's the retina along with the choroid and underlying sclera.

- --First, while I realize the neurosensory retina looks wonky, that's not the point of the slide.
- --The choroid is filled with innumerable inflammatory cells.
- --Two focal aggregates of inflammatory cells can be seen **here** and **here**. Drill down on these aggregates—exactly where are they located? Note what's running across their tops—a thin, densely pigmented line. This is the RPE.

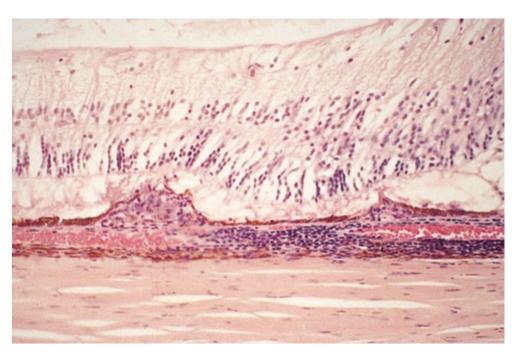




Hopefully you recognize what we're looking at here—it's the retina along with the choroid and underlying sclera.

- --First, while I realize the neurosensory retina looks wonky, that's not the point of the slide.
- --The choroid is filled with innumerable inflammatory cells.
- --Two focal aggregates of inflammatory cells can be seen **here** and **here**. Drill down on these aggregates—exactly where are they located? Note what's running across their tops—a thin, densely pigmented line. This is the RPE. And because the RPE is separated from the underlying choroid by eponym membrane, we can't be sure if the aggregates are below or just above it.

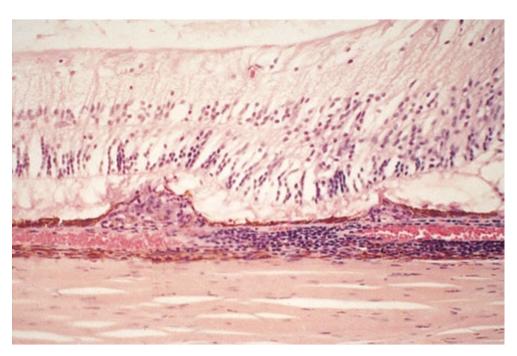




Hopefully you recognize what we're looking at here—it's the retina along with the choroid and underlying sclera.

- --First, while I realize the neurosensory retina looks wonky, that's not the point of the slide.
- --The choroid is filled with innumerable inflammatory cells.
- --Two focal aggregates of inflammatory cells can be seen **here** and **here**. Drill down on these aggregates—exactly where are they located? Note what's running across their tops—a thin, densely pigmented line. This is the RPE. And because the RPE is separated from the underlying choroid by Bruch's membrane, we can't be sure if the aggregates are *below* Bruch's or just *above* it.





What's the finding?*

Hopefully you recognize what we're looking at here—it's the retina along with the choroid and underlying sclera.

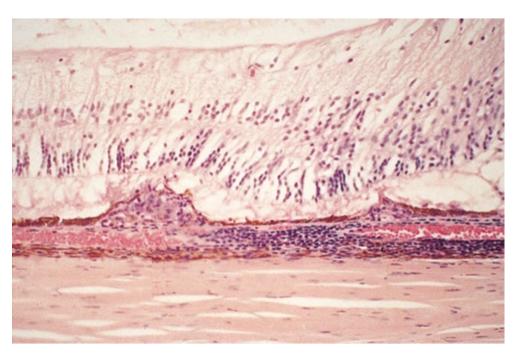
What's going on here—what are we supposed to take note of?

- --First, while I realize the neurosensory retina looks wonky, that's not the point of the slide.
- --The choroid is filled with innumerable inflammatory cells.
- --Two focal aggregates of inflammatory cells can be seen **here** and **here**. Drill down on these aggregates—exactly where are they located? Note what's running across their tops—a thin, densely pigmented line. This is the RPE. And because the RPE is separated from the underlying choroid by Bruch's membrane, we can't be sure if the aggregates are *below* Bruch's or just *above* it.

Focal aggregates of inflammatory cells just beneath the RPE + widespread choroidal inflammatory infiltration points to one entity:

^{*}Note that we're looking for the name of the aggregates, not the name of the underlying condition causing them.





What's the finding?

Hopefully you recognize what we're looking at here—it's the retina along with the choroid and underlying sclera.

What's going on here—what are we supposed to take note of?

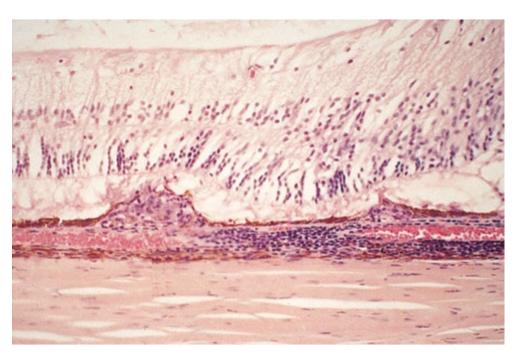
- --First, while I realize the neurosensory retina looks wonky, that's not the point of the slide.
- --The choroid is filled with innumerable inflammatory cells.
- --Two focal aggregates of inflammatory cells can be seen **here** and **here**. Drill down on these aggregates—exactly where are they located? Note what's running across their tops—a thin, densely pigmented line. This is the RPE. And because the RPE is separated from the underlying choroid by Bruch's membrane, we can't be sure if the aggregates are *below* Bruch's or just *above* it.

Focal aggregates of inflammatory cells just beneath the RPE + widespread choroidal inflammatory infiltration points to one entity:

Dalen-Fuchs nodules are inflammatory-cell aggregates between

two structures





What's the finding?

Hopefully you recognize what we're looking at here—it's the retina along with the choroid and underlying sclera.

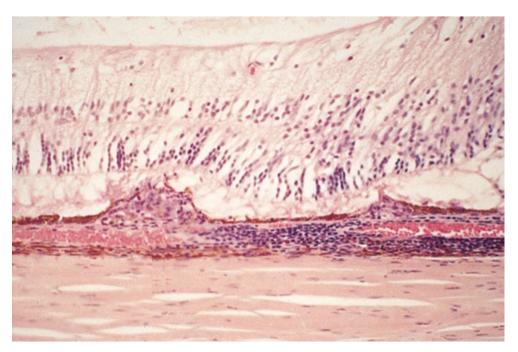
What's going on here—what are we supposed to take note of?

- --First, while I realize the neurosensory retina looks wonky, that's not the point of the slide.
- --The choroid is filled with innumerable inflammatory cells.
- --Two focal aggregates of inflammatory cells can be seen **here** and **here**. Drill down on these aggregates—exactly where are they located? Note what's running across their tops—a thin, densely pigmented line. This is the RPE. And because the RPE is separated from the underlying choroid by Bruch's membrane, we can't be sure if the aggregates are *below* Bruch's or just *above* it.

Focal aggregates of inflammatory cells just beneath the RPE + widespread choroidal inflammatory infiltration points to one entity:

Dalen-Fuchs nodules are inflammatory-cell aggregates between the RPE and Bruch's.





What's the finding?

Hopefully you recognize what we're looking at here—it's the retina along with the choroid and underlying sclera.

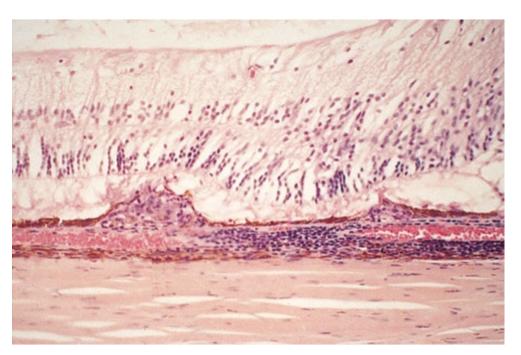
What's going on here—what are we supposed to take note of?

- --First, while I realize the neurosensory retina looks wonky, that's not the point of the slide.
- --The choroid is filled with innumerable inflammatory cells.
- --Two focal aggregates of inflammatory cells can be seen **here** and **here**. Drill down on these aggregates—exactly where are they located? Note what's running across their tops—a thin, densely pigmented line. This is the RPE. And because the RPE is separated from the underlying choroid by Bruch's membrane, we can't be sure if the aggregates are *below* Bruch's or just *above* it.

Focal aggregates of inflammatory cells just beneath the RPE + widespread choroidal inflammatory infiltration points to one entity:

Dalen-Fuchs nodules are inflammatory-cell aggregates between the RPE and Bruch's. They are most strongly associated with two conditions:





What's the finding?

Hopefully you recognize what we're looking at here—it's the retina along with the choroid and underlying sclera.

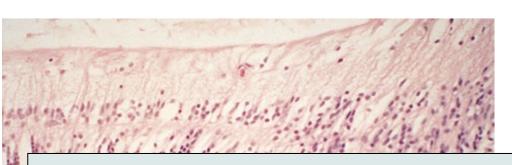
What's going on here—what are we supposed to take note of?

- --First, while I realize the neurosensory retina looks wonky, that's not the point of the slide.
- --The choroid is filled with innumerable inflammatory cells.
- --Two focal aggregates of inflammatory cells can be seen **here** and **here**. Drill down on these aggregates—exactly where are they located? Note what's running across their tops—a thin, densely pigmented line. This is the RPE. And because the RPE is separated from the underlying choroid by Bruch's membrane, we can't be sure if the aggregates are *below* Bruch's or just *above* it.

Focal aggregates of inflammatory cells just beneath the RPE + widespread choroidal inflammatory infiltration points to one entity:

Dalen-Fuchs nodules are inflammatory-cell aggregates between the RPE and Bruch's. They are most strongly associated with two conditions: Vogt-Koyanagi-Harada syndrome (VKH) and sympathetic ophthalmia (SO).



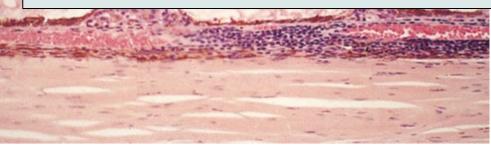


Hopefully you recognize what we're looking at here—it's the retina along with the choroid and underlying sclera.

What's going on here—what are we supposed to take note of?

- --First, while I realize the neurosensory retina looks wonky, that's not the point of the slide.
- --The choroid is filled with innumerable inflammatory cells.
- --Two focal aggregates of inflammatory cells can be seen **here** and **here**. Drill down on these

For more on Dalen-Fuchs nodules, see slide-set U6

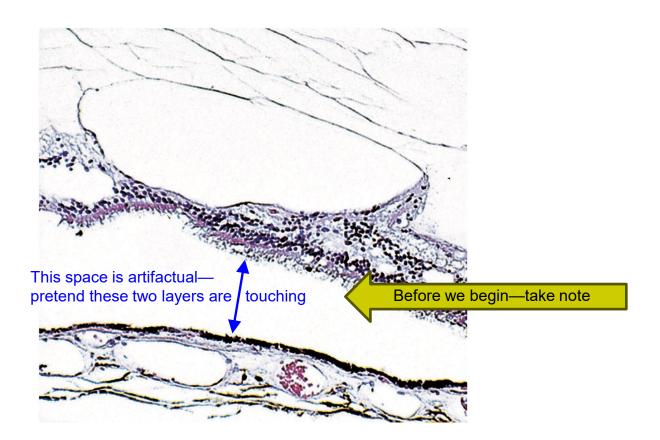


What's the finding?

And because the RPE is separated from the underlying choroid by Bruch's membrane, we can't be sure if the aggregates are *below* Bruch's or just *above* it.

Focal aggregates of inflammatory cells just beneath the RPE + widespread choroidal inflammatory infiltration points to one entity:

Dalen-Fuchs nodules are inflammatory-cell aggregates between the RPE and Bruch's. They are most strongly associated with two conditions: Vogt-Koyanagi-Harada syndrome (VKH) and sympathetic ophthalmia (SO).





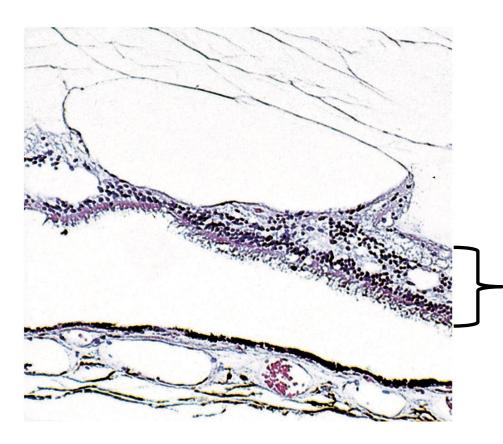




We're looking at four basic structures here: --[the one marked by the bracket]

- --?
- --?
- --2

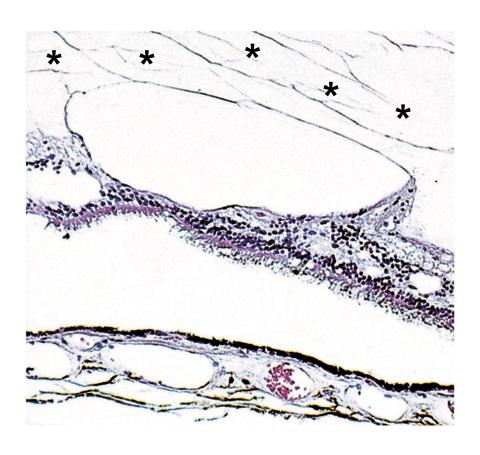




We're looking at four basic structures here:

- --The neurosensory (NS) retina
- -- 7
- --?
- __2





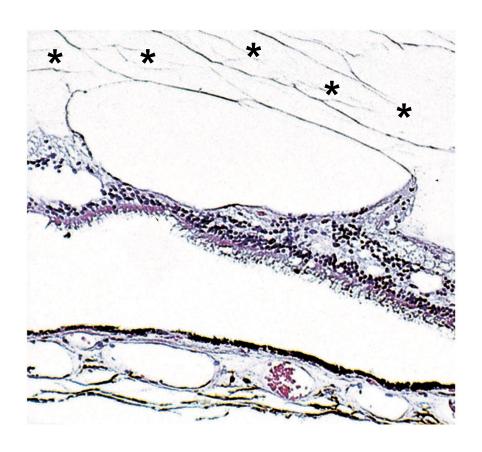
We're looking at four basic structures here:

- --The neurosensory (NS) retina
- --[the asterisks]

-- 2

--?





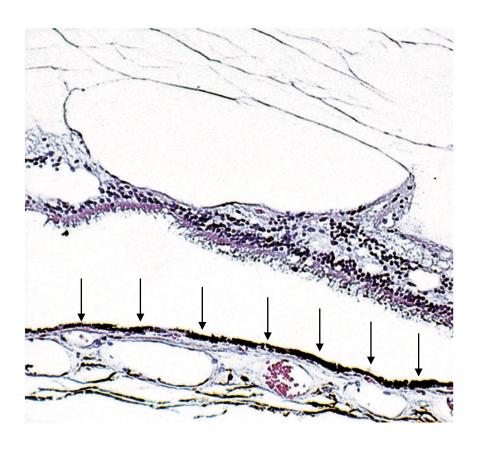
We're looking at four basic structures here:

- --The neurosensory (NS) retina
- --The vitreous

-- 2

-- ?



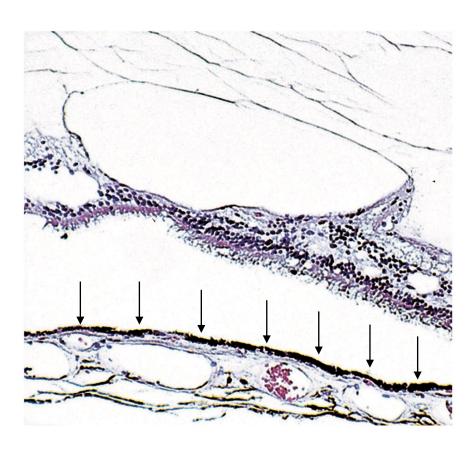


We're looking at four basic structures here:

- --The neurosensory (NS) retina
- --The vitreous
- --[the arrows]

-- 7



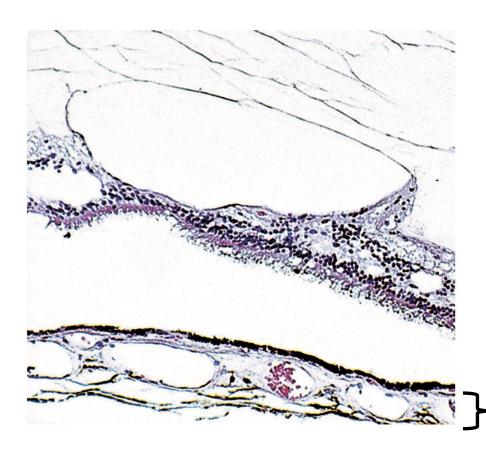


We're looking at four basic structures here:

- --The neurosensory (NS) retina
- --The vitreous
- --The RPE

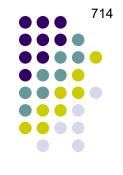
-- 7

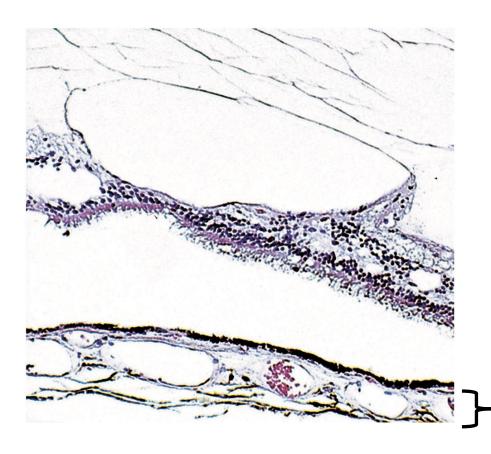




We're looking at four basic structures here:

- --The neurosensory (NS) retina
- --The vitreous
- --The RPE
- --[the bracket]





We're looking at four basic structures here:

- --The neurosensory (NS) retina
- --The vitreous
- --The RPE
- --The choroid





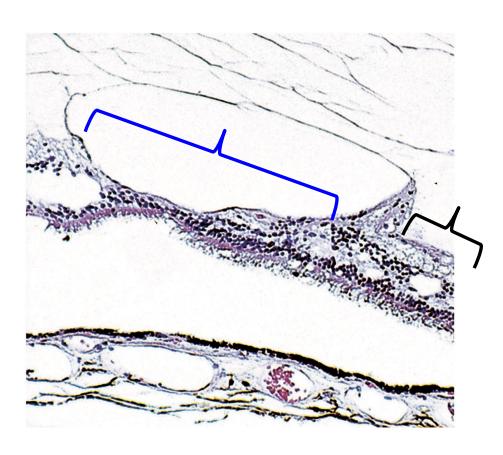
We're looking at four basic structures here:

- --The neurosensory (NS) retina
- --The vitreous
- --The RPE
- --The choroid

As for what's going on here, let's take note of the following:

--If we take the NS retina from **here** to **here** as being normal-ish...





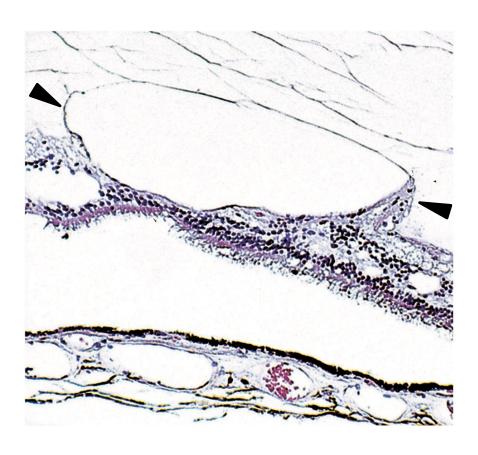
We're looking at four basic structures here:

- --The neurosensory (NS) retina
- --The vitreous
- --The RPE
- --The choroid

As for what's going on here, let's take note of the following:

--If we take the NS retina from **here** to **here** as being normal-ish...we can see that from here to here it is thin-to-very-thin.





We're looking at four basic structures here:

- --The neurosensory (NS) retina
- --The vitreous
- --The RPE
- --The choroid

As for what's going on here, let's take note of the following:

- --If we take the NS retina from **here** to **here** as being normal-ish...we can see that from here to here it is thin-to-very-thin.
- --The vitreous is attached to the area of thinned retina at its edges (*arrowheads*).





We're looking at four basic structures here:

- -- The neurosensory (NS) retina
- --The vitreous
- --The RPE
- --The choroid

As for what's going on here, let's take note of the following:

- --If we take the NS retina from **here** to **here** as being normal-ish...we can see that from here to here it is thin-to-very-thin.
- --The vitreous is attached to the area of thinned retina at its edges (*arrowheads*).
- --The vitreous directly overlying the thinned area is devoid of normal strands (*asterisks*).





What's the diagnosis?

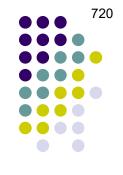
We're looking at four basic structures here:

- --The neurosensory (NS) retina
- --The vitreous
- --The RPE
- --The choroid

As for what's going on here, let's take note of the following:

- --If we take the NS retina from **here** to **here** as being normal-ish...we can see that from here to here it is thin-to-very-thin.
- --The vitreous is attached to the area of thinned retina at its edges (*arrowheads*).
- --The vitreous directly overlying the thinned area is devoid of normal strands (*asterisks*).

Focal retinal thinning attached to the V at its edges and associated with an overlying pocket of empty V is only one thing:





What's the diagnosis?

We're looking at four basic structures here:

- --The neurosensory (NS) retina
- -- The vitreous
- --The RPE
- --The choroid

As for what's going on here, let's take note of the following:

- --If we take the NS retina from **here** to **here** as being normal-ish...we can see that from here to here it is thin-to-very-thin.
- --The vitreous is attached to the area of thinned retina at its edges (*arrowheads*).
- --The vitreous directly overlying the thinned area is devoid of normal strands (*asterisks*).

Focal retinal thinning attached to the V at its edges and associated with an overlying pocket of empty V is only one thing:

Lattice degeneration is a common condition that may be associated with

three words





What's the diagnosis?

We're looking at four basic structures here:

- --The neurosensory (NS) retina
- -- The vitreous
- --The RPE
- --The choroid

As for what's going on here, let's take note of the following:

- --If we take the NS retina from **here** to **here** as being normal-ish...we can see that from here to here it is thin-to-very-thin.
- --The vitreous is attached to the area of thinned retina at its edges (*arrowheads*).
- --The vitreous directly overlying the thinned area is devoid of normal strands (*asterisks*).

Focal retinal thinning attached to the V at its edges and associated with an overlying pocket of empty V is only one thing:

Lattice degeneration is a common condition that may be associated with rhegmatogenous RD.





What's the diagnosis?

We're looking at four basic structures here:

- --The neurosensory (NS) retina
- -- The vitreous
- --The RPE
- --The choroid

As for what's going on here, let's take note of the following:

- --If we take the NS retina from **here** to **here** as being normal-ish...we can see that from here to here it is thin-to-very-thin.
- --The vitreous is attached to the area of thinned retina at its edges (*arrowheads*).
- --The vitreous directly overlying the thinned area is devoid of normal strands (*asterisks*).

Focal retinal thinning attached to the V at its edges and associated with an overlying pocket of empty V is only one thing:

Lattice degeneration is a common condition that may be associated with rhegmatogenous RD. Its chief features are atrophy of the retina





What's the diagnosis?

We're looking at four basic structures here:

- --The neurosensory (NS) retina
- -- The vitreous
- --The RPE
- --The choroid

As for what's going on here, let's take note of the following:

- --If we take the NS retina from **here** to **here** as being normal-ish...we can see that from here to here it is thin-to-very-thin.
- --The vitreous is attached to the area of thinned retina at its edges (*arrowheads*).
- --The vitreous directly overlying the thinned area is devoid of normal strands (*asterisks*).

Focal retinal thinning attached to the V at its edges and associated with an overlying pocket of empty V is only one thing:

Lattice degeneration is a common condition that may be associated with rhegmatogenous RD. Its chief features are atrophy of the inner retina





What's the diagnosis?

We're looking at four basic structures here:

- --The neurosensory (NS) retina
- -- The vitreous
- --The RPE
- --The choroid

As for what's going on here, let's take note of the following:

- --If we take the NS retina from **here** to **here** as being normal-ish...we can see that from here to here it is thin-to-very-thin.
- --The vitreous is attached to the area of thinned retina at its edges (*arrowheads*).
- --The vitreous directly overlying the thinned area is devoid of normal strands (*asterisks*).

Focal retinal thinning attached to the V at its edges and associated with an overlying pocket of empty V is only one thing:

Lattice degeneration is a common condition that may be associated with rhegmatogenous RD. Its chief features are atrophy of the inner retina (including absence of the labb.)





What's the diagnosis?

We're looking at four basic structures here:

- --The neurosensory (NS) retina
- -- The vitreous
- --The RPE
- --The choroid

As for what's going on here, let's take note of the following:

- --If we take the NS retina from **here** to **here** as being normal-ish...we can see that from here to here it is thin-to-very-thin.
- --The vitreous is attached to the area of thinned retina at its edges (*arrowheads*).
- --The vitreous directly overlying the thinned area is devoid of normal strands (*asterisks*).

Focal retinal thinning attached to the V at its edges and associated with an overlying pocket of empty V is only one thing:

Lattice degeneration is a common condition that may be associated with rhegmatogenous RD. Its chief features are atrophy of the inner retina (including absence of the ILM)





What's the diagnosis?

We're looking at four basic structures here:

- --The neurosensory (NS) retina
- -- The vitreous
- --The RPE
- --The choroid

As for what's going on here, let's take note of the following:

- --If we take the NS retina from **here** to **here** as being normal-ish...we can see that from here to here it is thin-to-very-thin.
- --The vitreous is attached to the area of thinned retina at its edges (*arrowheads*).
- --The vitreous directly overlying the thinned area is devoid of normal strands (*asterisks*).

Focal retinal thinning attached to the V at its edges and associated with an overlying pocket of empty V is only one thing:

Lattice degeneration is a common condition that may be associated with rhegmatogenous RD. Its chief features are atrophy of the inner retina (including absence of the ILM); an overlying pocket of vitreous





What's the diagnosis?

We're looking at four basic structures here:

- --The neurosensory (NS) retina
- -- The vitreous
- --The RPE
- -- The choroid

As for what's going on here, let's take note of the following:

- --If we take the NS retina from **here** to **here** as being normal-ish...we can see that from here to here it is thin-to-very-thin.
- --The vitreous is attached to the area of thinned retina at its edges (*arrowheads*).
- --The vitreous directly overlying the thinned area is devoid of normal strands (*asterisks*).

Focal retinal thinning attached to the V at its edges and associated with an overlying pocket of empty V is only one thing:

Lattice degeneration is a common condition that may be associated with rhegmatogenous RD. Its chief features are atrophy of the inner retina (including absence of the ILM); an overlying pocket of liquified vitreous





What's the diagnosis?

We're looking at four basic structures here:

- --The neurosensory (NS) retina
- --The vitreous
- --The RPE
- -- The choroid

As for what's going on here, let's take note of the following:

- --If we take the NS retina from **here** to **here** as being normal-ish...we can see that from here to here it is thin-to-very-thin.
- --The vitreous is attached to the area of thinned retina at its edges (*arrowheads*).
- --The vitreous directly overlying the thinned area is devoid of normal strands (*asterisks*).

Focal retinal thinning attached to the V at its edges and associated with an overlying pocket of empty V is only one thing:

Lattice degeneration is a common condition that may be associated with rhegmatogenous RD. Its chief features are atrophy of the inner retina (including absence of the ILM); an overlying pocket of liquified vitreous; and the weak adherence of vitreous at the outer boundary of the area.





What's the diagnosis?

We're looking at four basic structures here:

- --The neurosensory (NS) retina
- -- The vitreous
- --The RPE
- --The choroid

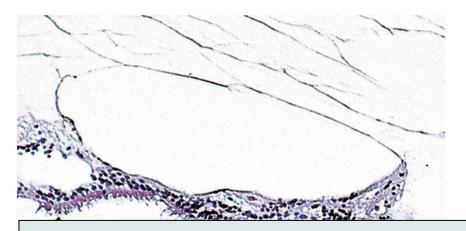
As for what's going on here, let's take note of the following:

- --If we take the NS retina from **here** to **here** as being normal-ish...we can see that from here to here it is thin-to-very-thin.
- --The vitreous is attached to the area of thinned retina at its edges (*arrowheads*).
- --The vitreous directly overlying the thinned area is devoid of normal strands (*asterisks*).

Focal retinal thinning attached to the V at its edges and associated with an overlying pocket of empty V is only one thing:

Lattice degeneration is a common condition that may be associated with rhegmatogenous RD. Its chief features are atrophy of the inner retina (including absence of the ILM); an overlying pocket of liquified vitreous; and the firm adherence of vitreous at the outer boundary of the area.





We're looking at four basic structures here:

- --The neurosensory (NS) retina
- -- The vitreous
- --The RPE
- --The choroid

As for what's going on here, let's take note of the following:

If we take the NS retina from hare to hare as

For more on lattice degeneration, see slide-set R36



What's the diagnosis?

- --The vitreous is attached to the area of thinned retina at its edges (*arrowheads*).
- --The vitreous directly overlying the thinned area is devoid of normal strands (*asterisks*).

Focal retinal thinning attached to the V at its edges and associated with an overlying pocket of empty V is only one thing:

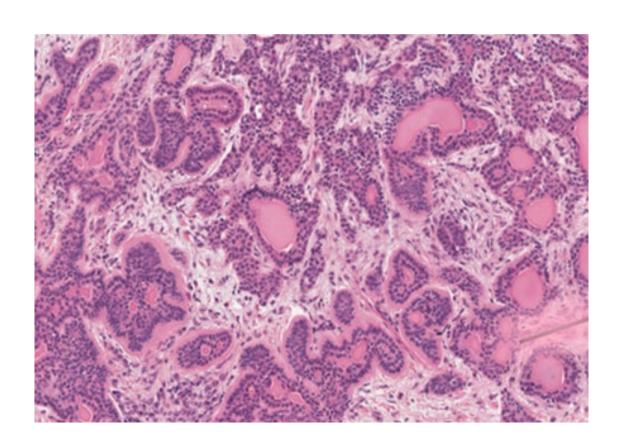
Lattice degeneration is a common condition that may be associated with rhegmatogenous RD. Its chief features are atrophy of the inner retina (including absence of the ILM); an overlying pocket of liquified vitreous; and the firm adherence of vitreous at the outer boundary of the area.



Images Consisting of Wall-to-Wall Cells

These images are challenging because they lack context—no up or down; no natural edges—thus making it difficult to know what the tissue is. That being said, if you're expected to work with such an image, *it will contain a field mark that gives its identity away*. Know the mark, know the diagnosis!

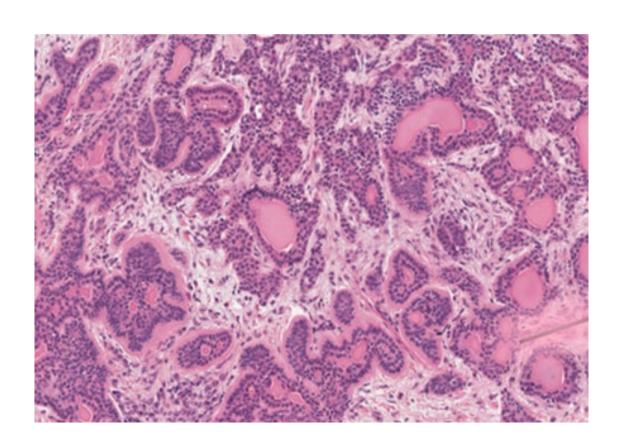




Note:

--The many glandular/tubular structures. For us eyedocs this suggests one tissue:

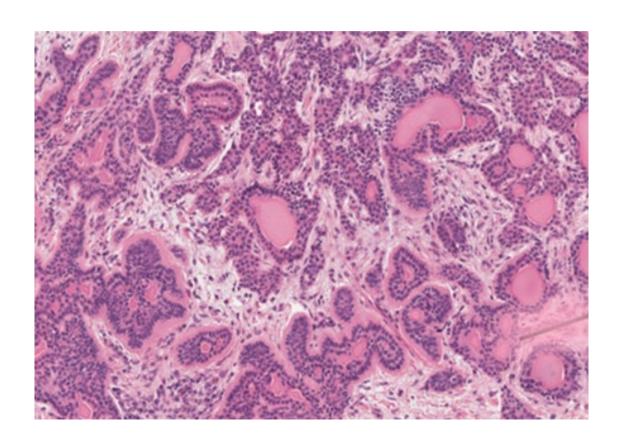




Note:

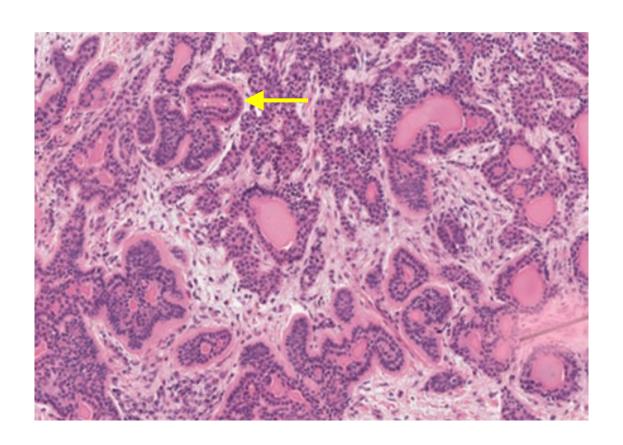
--The many glandular/tubular structures. For us eyedocs this suggests one tissue: The main lacrimal gland.





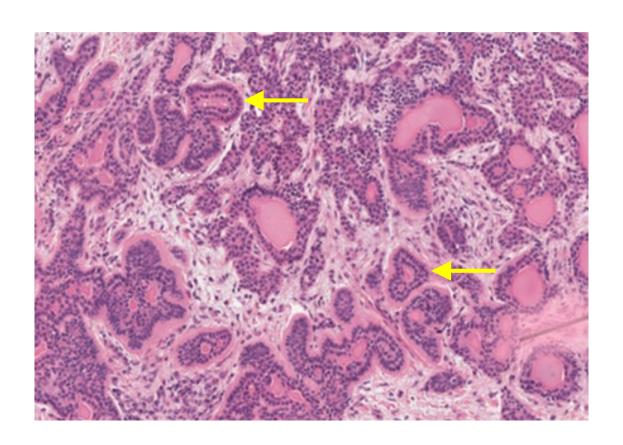
- --The many glandular/tubular structures. For us eyedocs this suggests one tissue: The main lacrimal gland.
- --The gland/tubule walls consist of two epi layers. This arrangement is consistently orderly throughout





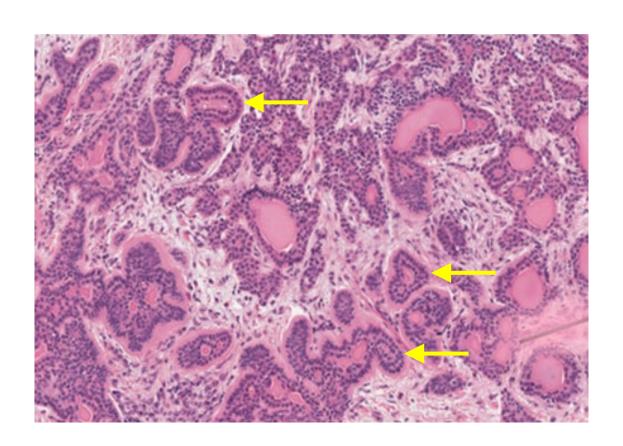
- --The many glandular/tubular structures. For us eyedocs this suggests one tissue: The main lacrimal gland.
- --The gland/tubule walls consist of two epi layers. This arrangement is consistently orderly throughout look here





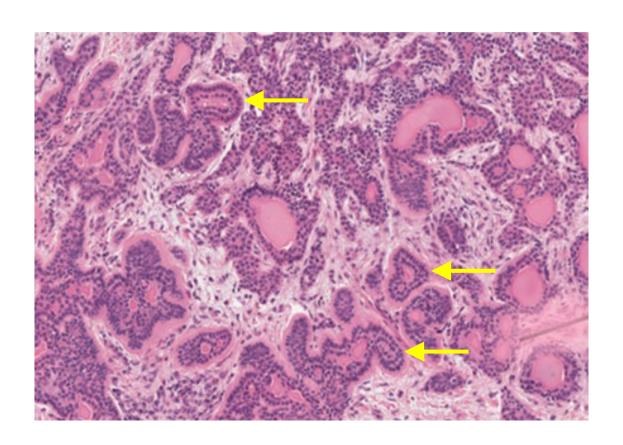
- --The many glandular/tubular structures. For us eyedocs this suggests one tissue: The main lacrimal gland.
- --The gland/tubule walls consist of two epi layers. This arrangement is consistently orderly throughout look here, here





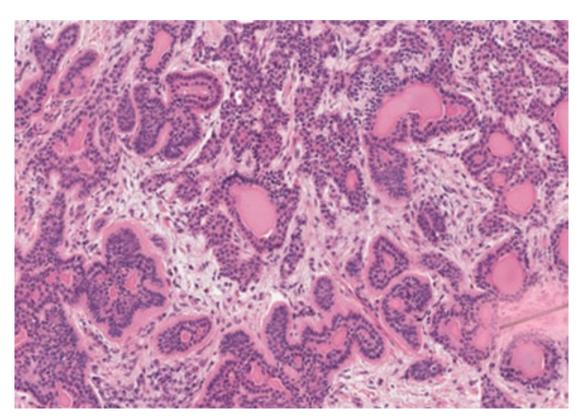
- --The many glandular/tubular structures. For us eyedocs this suggests one tissue: The main lacrimal gland.
- --The gland/tubule walls consist of two epi layers. This arrangement is consistently orderly throughout look here, here, and here, for example.





- --The many glandular/tubular structures. For us eyedocs this suggests one tissue: The main lacrimal gland.
- --The gland/tubule walls consist of two epi layers. This arrangement is consistently orderly throughout look here, here, and here, for example. In short, there's nothing here to suggest the chaotic, uncontrolled growth of a malignancy.

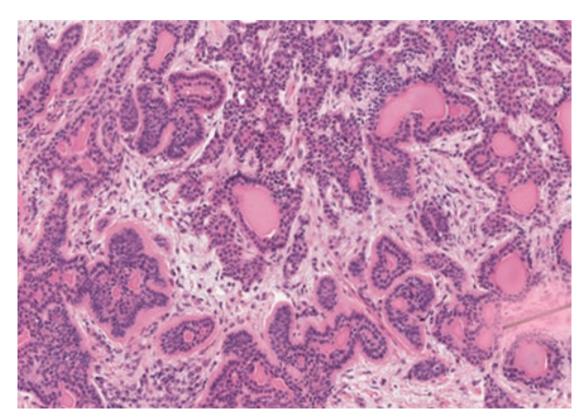




What's the diagnosis?

- --The many glandular/tubular structures. For us eyedocs this suggests one tissue: The main lacrimal gland.
- --The gland/tubule walls consist of two epi layers. This arrangement is consistently orderly throughout look here, here, and here, for example. In short, there's nothing here to suggest the chaotic, uncontrolled growth of a malignancy.





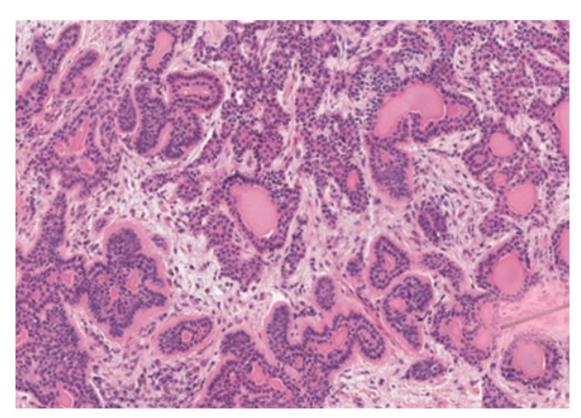
Note:

- --The many glandular/tubular structures. For us eyedocs this suggests one tissue: The main lacrimal gland.
- --The gland/tubule walls consist of two epi layers. This arrangement is consistently orderly throughout look here, here, and here, for example. In short, there's nothing here to suggest the chaotic, uncontrolled growth of a malignancy.

What's the diagnosis?

Pleomorphic adenoma, the common epithelial tumor of the lacrimal gland





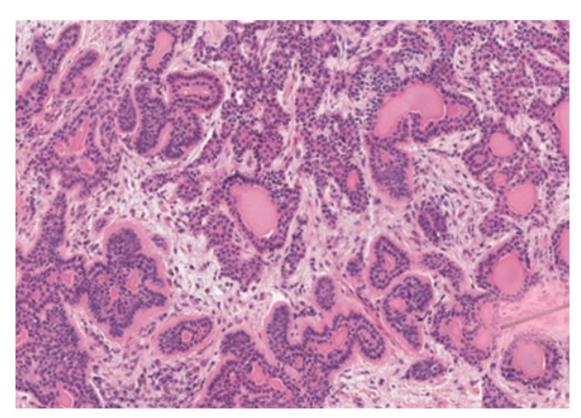
Note:

- --The many glandular/tubular structures. For us eyedocs this suggests one tissue: The main lacrimal gland.
- --The gland/tubule walls consist of two epi layers. This arrangement is consistently orderly throughout look here, here, and here, for example. In short, there's nothing here to suggest the chaotic, uncontrolled growth of a malignancy.

What's the diagnosis?

Pleomorphic adenoma, the most common epithelial tumor of the lacrimal gland





Note:

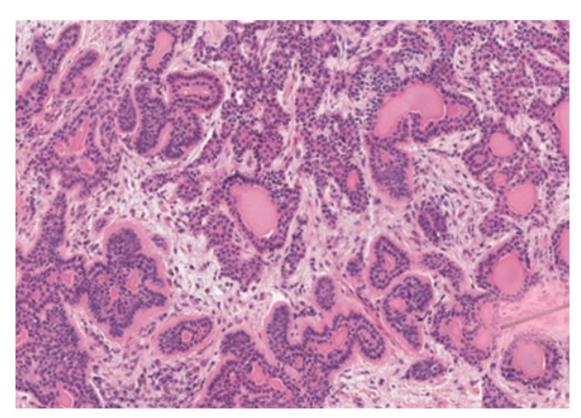
- --The many glandular/tubular structures. For us eyedocs this suggests one tissue: The main lacrimal gland.
- --The gland/tubule walls consist of two epi layers. This arrangement is consistently orderly throughout look here, here, and here, for example. In short, there's nothing here to suggest the chaotic, uncontrolled growth of a malignancy.

What's the diagnosis?

Pleomorphic adenoma, the most common epithelial tumor of the lacrimal gland. It is







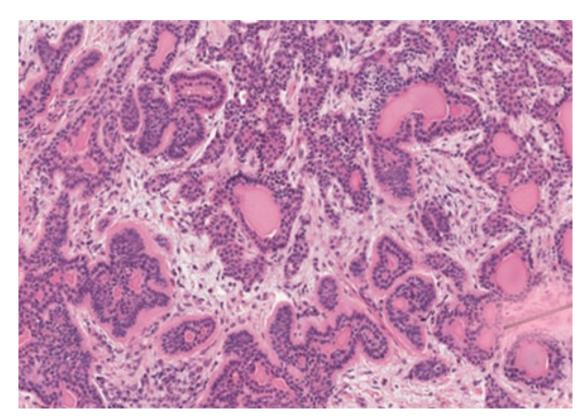
Note:

- --The many glandular/tubular structures. For us eyedocs this suggests one tissue: The main lacrimal gland.
- --The gland/tubule walls consist of two epi layers. This arrangement is consistently orderly throughout look here, here, and here, for example. In short, there's nothing here to suggest the chaotic, uncontrolled growth of a malignancy.

What's the diagnosis?

Pleomorphic adenoma, the most common epithelial tumor of the lacrimal gland. It is painless.





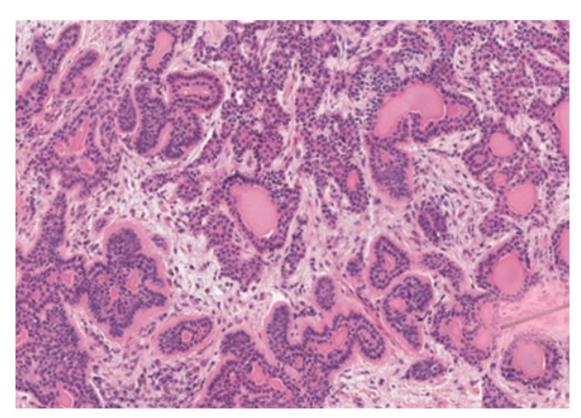
Note:

- --The many glandular/tubular structures. For us eyedocs this suggests one tissue: The main lacrimal gland.
- --The gland/tubule walls consist of two epi layers. This arrangement is consistently orderly throughout look here, here, and here, for example. In short, there's nothing here to suggest the chaotic, uncontrolled growth of a malignancy.

What's the diagnosis?

Pleomorphic adenoma, the most common epithelial tumor of the lacrimal gland. It is painless. Slightly more common in Mrs





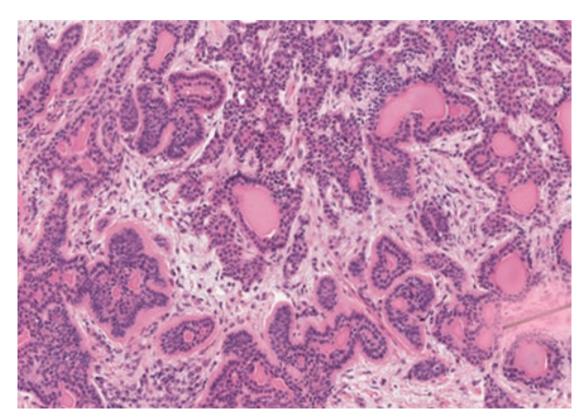
Note:

- --The many glandular/tubular structures. For us eyedocs this suggests one tissue: The main lacrimal gland.
- --The gland/tubule walls consist of two epi layers. This arrangement is consistently orderly throughout look here, here, and here, for example. In short, there's nothing here to suggest the chaotic, uncontrolled growth of a malignancy.

What's the diagnosis?

Pleomorphic adenoma, the most common epithelial tumor of the lacrimal gland. It is painless. Slightly more common in men.





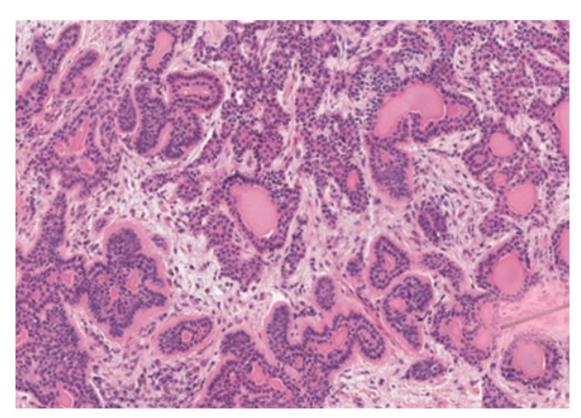
Note:

- --The many glandular/tubular structures. For us eyedocs this suggests one tissue: The main lacrimal gland.
- --The gland/tubule walls consist of two epi layers. This arrangement is consistently orderly throughout look here, here, and here, for example. In short, there's nothing here to suggest the chaotic, uncontrolled growth of a malignancy.

What's the diagnosis?

Pleomorphic adenoma, the most common epithelial tumor of the lacrimal gland. It is painless. Slightly more common in men. Presents in the decade.





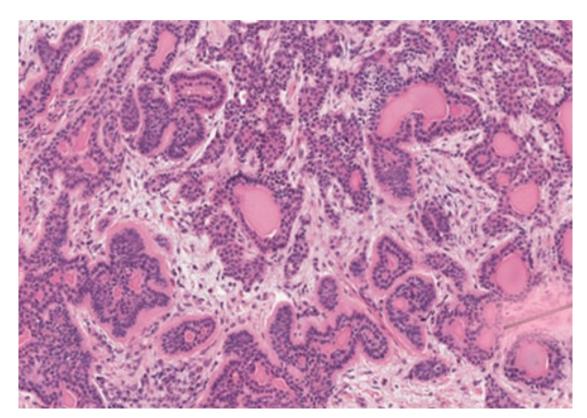
Note: --The

- --The many glandular/tubular structures. For us eyedocs this suggests one tissue: The main lacrimal gland.
- --The gland/tubule walls consist of two epi layers. This arrangement is consistently orderly throughout look here, here, and here, for example. In short, there's nothing here to suggest the chaotic, uncontrolled growth of a malignancy.

What's the diagnosis?

Pleomorphic adenoma, the most common epithelial tumor of the lacrimal gland. It is painless. Slightly more common in men. Presents in the 4^{th} - 5^{th} decade.





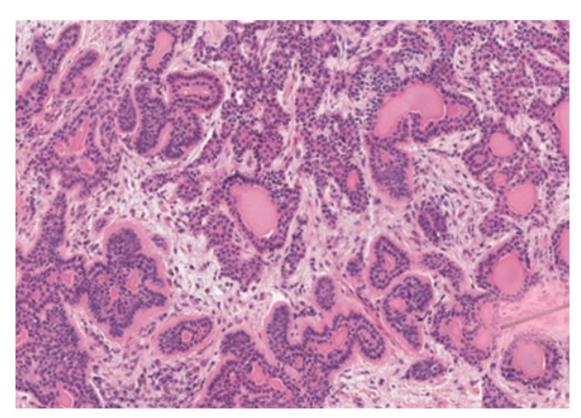
Note:

- --The many glandular/tubular structures. For us eyedocs this suggests one tissue: The main lacrimal gland.
- --The gland/tubule walls consist of two epi layers. This arrangement is consistently orderly throughout look here, here, and here, for example. In short, there's nothing here to suggest the chaotic, uncontrolled growth of a malignancy.

What's the diagnosis?

Pleomorphic adenoma, the most common epithelial tumor of the lacrimal gland. It is painless. Slightly more common in men. Presents in the 4th-5th decade. The tumor is encapsulated?





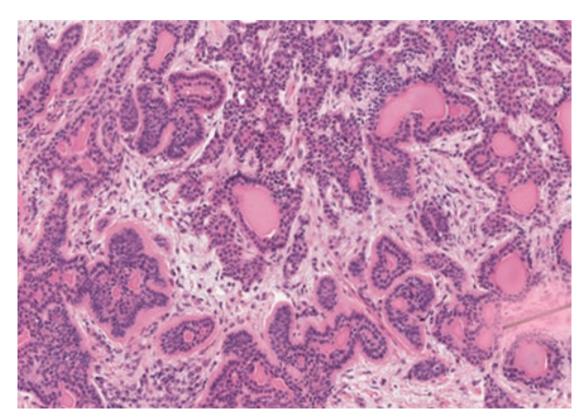
Note:

- --The many glandular/tubular structures. For us eyedocs this suggests one tissue: The main lacrimal gland.
- --The gland/tubule walls consist of two epi layers. This arrangement is consistently orderly throughout look here, here, and here, for example. In short, there's nothing here to suggest the chaotic, uncontrolled growth of a malignancy.

What's the diagnosis?

Pleomorphic adenoma, the most common epithelial tumor of the lacrimal gland. It is painless. Slightly more common in men. Presents in the 4th-5th decade. The tumor is pseudoencapsulated.





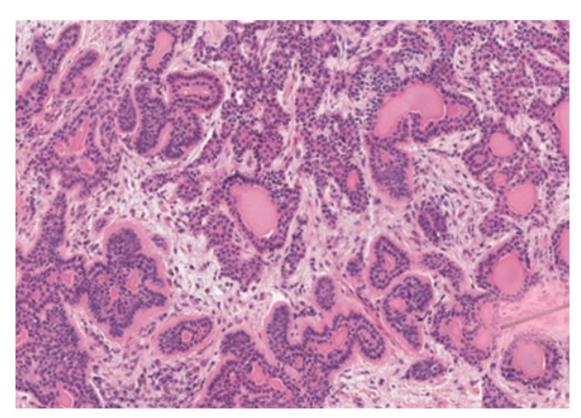
Note:

- --The many glandular/tubular structures. For us eyedocs this suggests one tissue: The main lacrimal gland.
- --The gland/tubule walls consist of two epi layers. This arrangement is consistently orderly throughout look here, here, and here, for example. In short, there's nothing here to suggest the chaotic, uncontrolled growth of a malignancy.

What's the diagnosis?

Pleomorphic adenoma, the most common epithelial tumor of the lacrimal gland. It is painless. Slightly more common in men. Presents in the 4th-5th decade. The tumor is pseudoencapsulated. Grows quickly v slowly





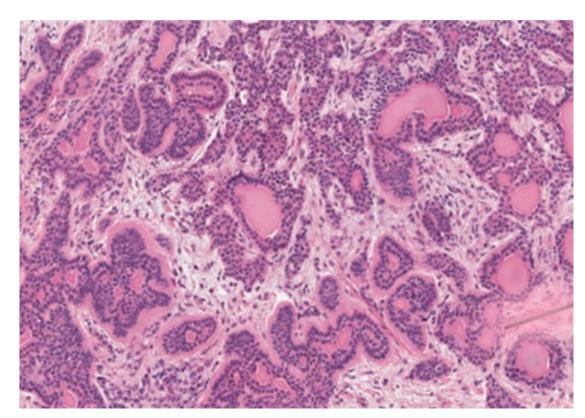
Note:

- --The many glandular/tubular structures. For us eyedocs this suggests one tissue: The main lacrimal gland.
- --The gland/tubule walls consist of two epi layers. This arrangement is consistently orderly throughout look here, here, and here, for example. In short, there's nothing here to suggest the chaotic, uncontrolled growth of a malignancy.

What's the diagnosis?

Pleomorphic adenoma, the most common epithelial tumor of the lacrimal gland. It is painless. Slightly more common in men. Presents in the 4th-5th decade. The tumor is pseudoencapsulated. Grows slowly.





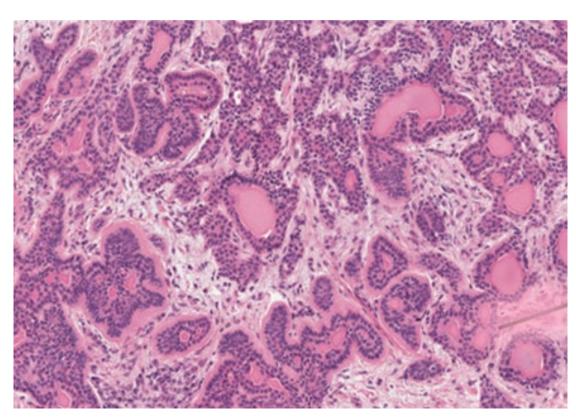
Note:

- --The many glandular/tubular structures. For us eyedocs this suggests one tissue: The main lacrimal gland.
- --The gland/tubule walls consist of two epi layers. This arrangement is consistently orderly throughout look here, here, and here, for example. In short, there's nothing here to suggest the chaotic, uncontrolled growth of a malignancy.

What's the diagnosis?

Pleomorphic adenoma, the most common epithelial tumor of the lacrimal gland. It is painless. Slightly more common in men. Presents in the 4th-5th decade. The tumor is pseudoencapsulated. Grows slowly. Progressive growth may adjacent bone but does not it.





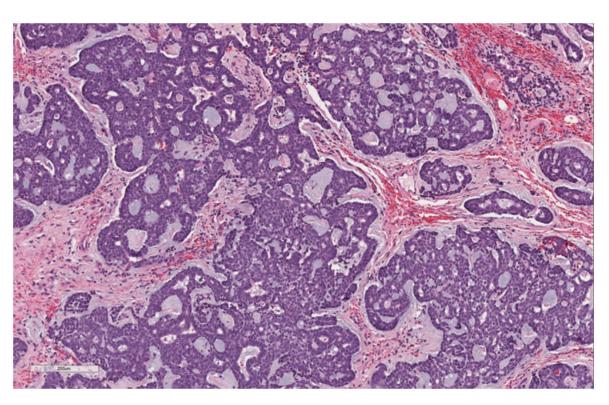
Note:

- --The many glandular/tubular structures. For us eyedocs this suggests one tissue: The main lacrimal gland.
- --The gland/tubule walls consist of two epi layers. This arrangement is consistently orderly throughout look here, here, and here, for example. In short, there's nothing here to suggest the chaotic, uncontrolled growth of a malignancy.

What's the diagnosis?

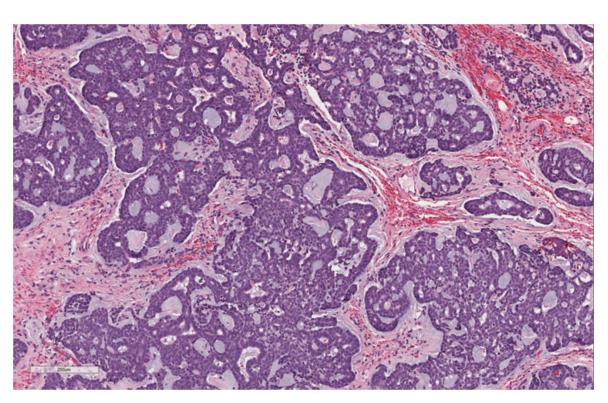
Pleomorphic adenoma, the most common epithelial tumor of the lacrimal gland. It is painless. Slightly more common in men. Presents in the 4th-5th decade. The tumor is pseudoencapsulated. Grows slowly. Progressive growth may excavate adjacent bone but does not erode it.





This tissue also forms glandular/tubular structures with lumen, and clearly represents the main lacrimal gland as well.

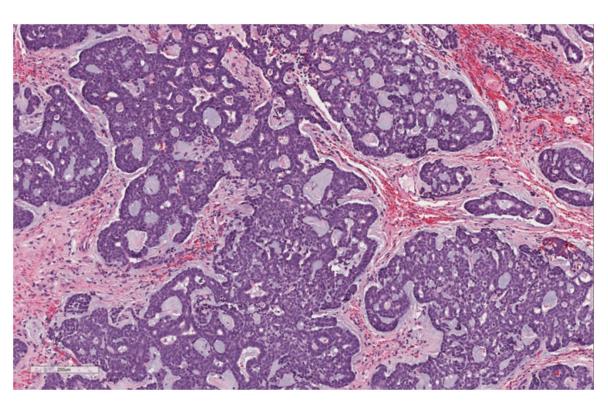




This tissue also forms glandular/tubular structures with lumen, and clearly represents the main lacrimal gland as well.

However, instead of two wellordered epi layers, the epi here is dense and chaotic.



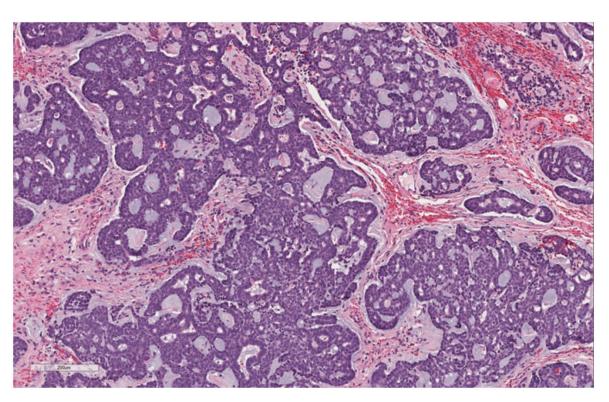


This tissue also forms glandular/tubular structures with lumen, and clearly represents the main lacrimal gland as well.

However, instead of two wellordered epi layers, the epi here is dense and chaotic. The multitude of small tubules is responsible for its appearance.

two-word classic description

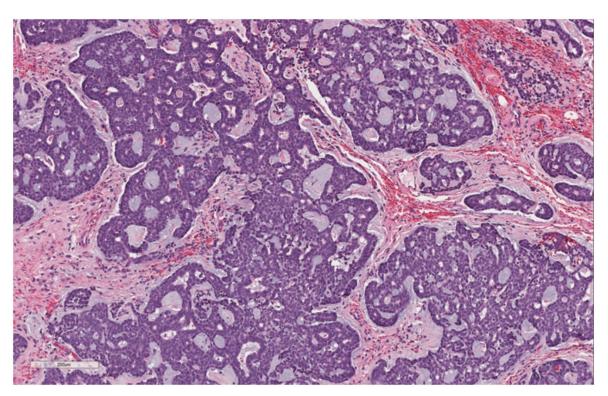




This tissue also forms glandular/tubular structures with lumen, and clearly represents the main lacrimal gland as well. However, instead of two well-ordered epi layers, the epi here is dense and chaotic. The multitude of small tubules is responsible for its

'Swiss cheese 'appearance.



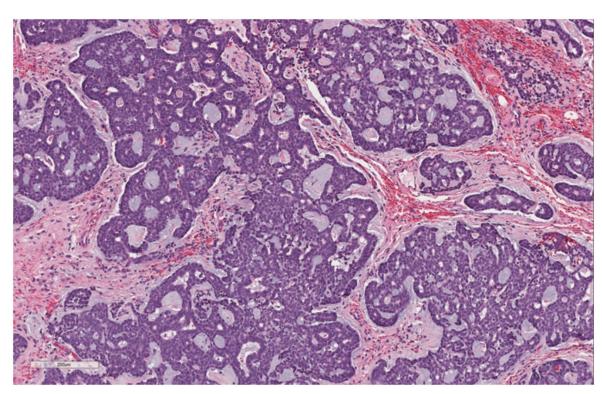


What's the diagnosis?

This tissue also forms glandular/tubular structures with lumen, and clearly represents the main lacrimal gland as well. However, instead of two well-ordered epi layers, the epi here is dense and chaotic. The multitude of small tubules is responsible for its

'Swiss cheese 'appearance.





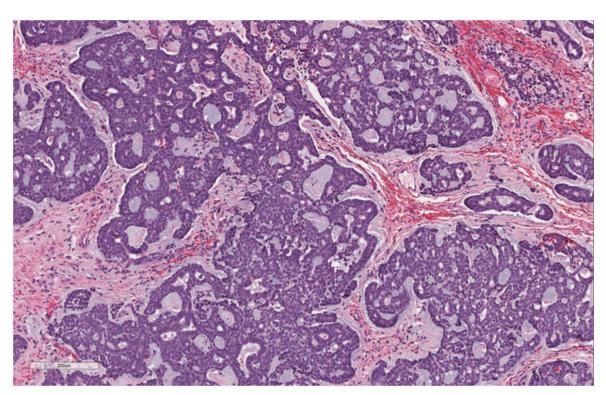
This tissue also forms glandular/tubular structures with lumen, and clearly represents the main lacrimal gland as well. However, instead of two well-ordered epi layers, the epi here is dense and chaotic. The multitude of small tubules is responsible for its 'Swiss cheese 'appearance.

What's the diagnosis?

Adenoid cystic carcinoma (ACC) is slightly more common in

M vs I



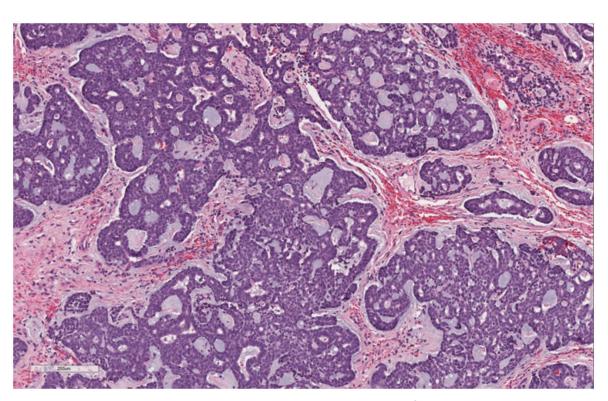


This tissue also forms glandular/tubular structures with lumen, and clearly represents the main lacrimal gland as well. However, instead of two well-ordered epi layers, the epi here is dense and chaotic. The multitude of small tubules is responsible for its 'Swiss cheese 'appearance.

What's the diagnosis?

Adenoid cystic carcinoma (ACC) is slightly more common in women



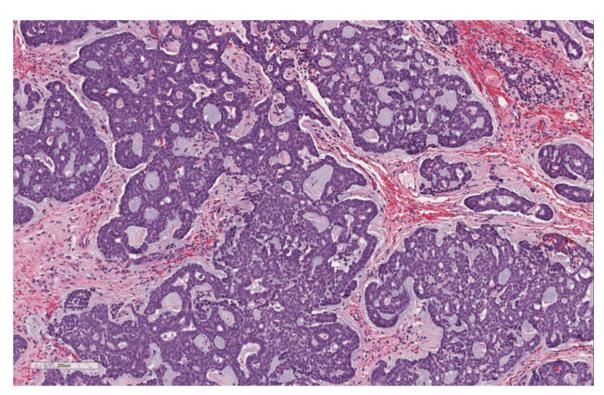


This tissue also forms glandular/tubular structures with lumen, and clearly represents the main lacrimal gland as well. However, instead of two well-ordered epi layers, the epi here is dense and chaotic. The multitude of small tubules is responsible for its 'Swiss cheese 'appearance.

What's the diagnosis?

Adenoid cystic carcinoma (ACC) is slightly more common in women. The median age at presentation is about # years



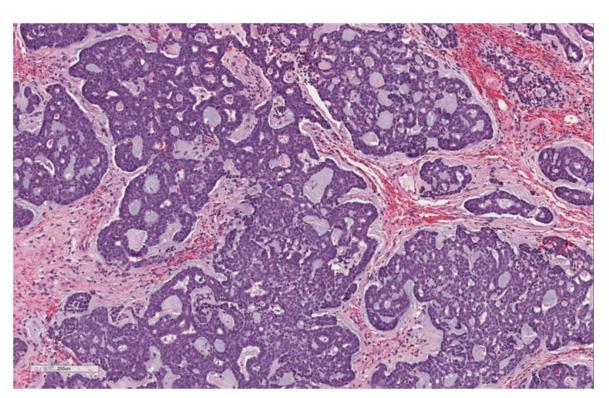


This tissue also forms glandular/tubular structures with lumen, and clearly represents the main lacrimal gland as well. However, instead of two well-ordered epi layers, the epi here is dense and chaotic. The multitude of small tubules is responsible for its 'Swiss cheese 'appearance.

What's the diagnosis?

Adenoid cystic carcinoma (ACC) is slightly more common in women. The median age at presentation is about 40 years.



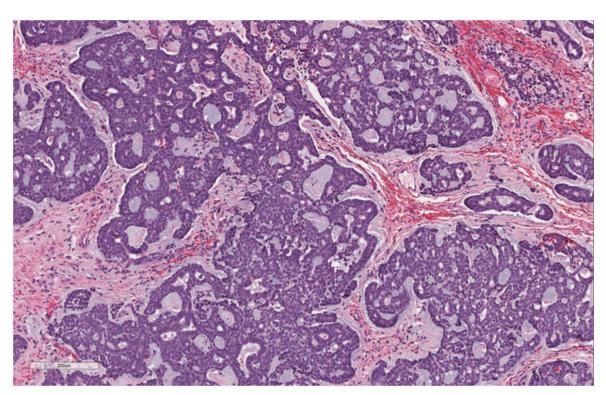


This tissue also forms glandular/tubular structures with lumen, and clearly represents the main lacrimal gland as well. However, instead of two well-ordered epi layers, the epi here is dense and chaotic. The multitude of small tubules is responsible for its 'Swiss cheese 'appearance.

What's the diagnosis?

Adenoid cystic carcinoma (ACC) is slightly more common in women. The median age at presentation is about 40 years. ACC has no pseudocapsule.



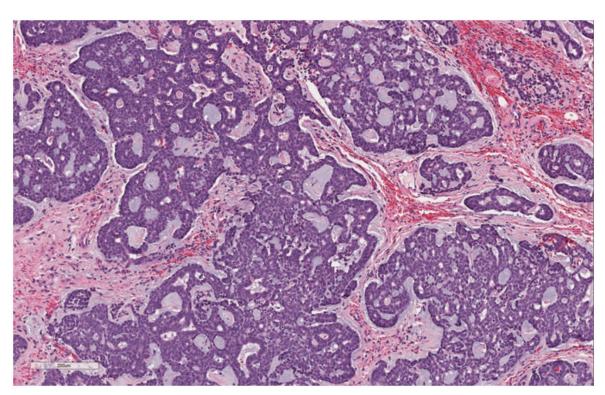


This tissue also forms glandular/tubular structures with lumen, and clearly represents the main lacrimal gland as well. However, instead of two well-ordered epi layers, the epi here is dense and chaotic. The multitude of small tubules is responsible for its 'Swiss cheese 'appearance.

What's the diagnosis?

Adenoid cystic carcinoma (ACC) is slightly more common in women. The median age at presentation is about 40 years. ACC has no pseudocapsule.



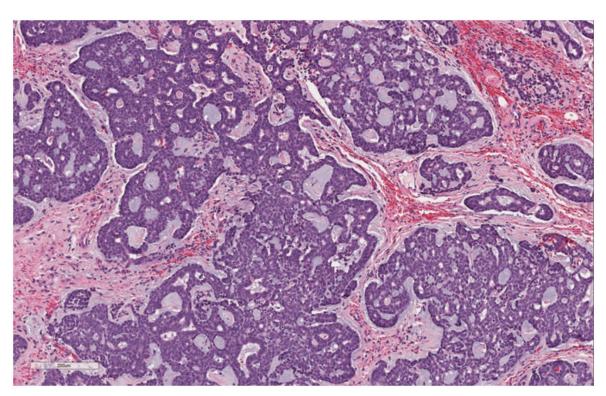


This tissue also forms glandular/tubular structures with lumen, and clearly represents the main lacrimal gland as well. However, instead of two well-ordered epi layers, the epi here is dense and chaotic. The multitude of small tubules is responsible for its 'Swiss cheese 'appearance.

What's the diagnosis?

Adenoid cystic carcinoma (ACC) is slightly more common in women. The median age at presentation is about 40 years. ACC has no pseudocapsule. Bone erosion is typical vs atypical vs atypical vs atypical vs atypical vs



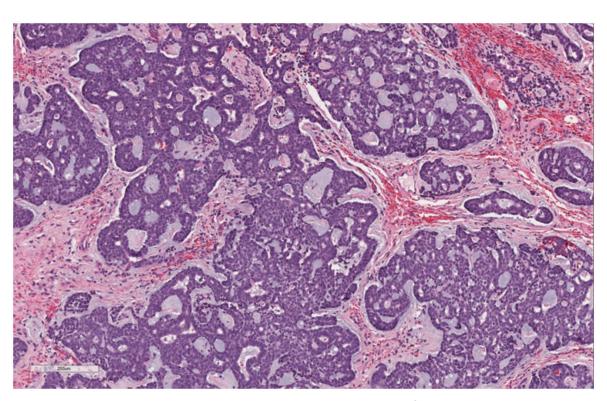


This tissue also forms glandular/tubular structures with lumen, and clearly represents the main lacrimal gland as well. However, instead of two well-ordered epi layers, the epi here is dense and chaotic. The multitude of small tubules is responsible for its 'Swiss cheese 'appearance.

What's the diagnosis?

Adenoid cystic carcinoma (ACC) is slightly more common in women. The median age at presentation is about 40 years. ACC has no pseudocapsule. Bone erosion is typical



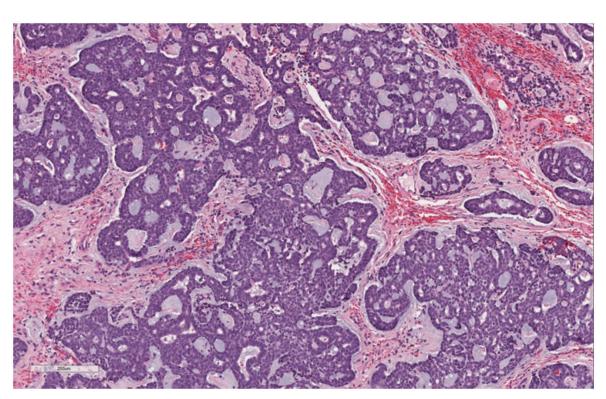


This tissue also forms glandular/tubular structures with lumen, and clearly represents the main lacrimal gland as well. However, instead of two well-ordered epi layers, the epi here is dense and chaotic. The multitude of small tubules is responsible for its 'Swiss cheese 'appearance.

What's the diagnosis?

Adenoid cystic carcinoma (ACC) is slightly more common in women. The median age at presentation is about 40 years. ACC has no pseudocapsule. Bone erosion is typical, and pain at presentation is common vs uncommon



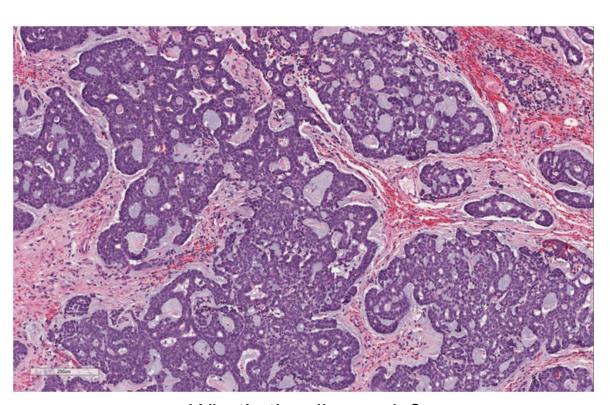


This tissue also forms glandular/tubular structures with lumen, and clearly represents the main lacrimal gland as well. However, instead of two well-ordered epi layers, the epi here is dense and chaotic. The multitude of small tubules is responsible for its 'Swiss cheese 'appearance.

What's the diagnosis?

Adenoid cystic carcinoma (ACC) is slightly more common in women. The median age at presentation is about 40 years. ACC has no pseudocapsule. Bone erosion is typical, and pain at presentation is the rule.



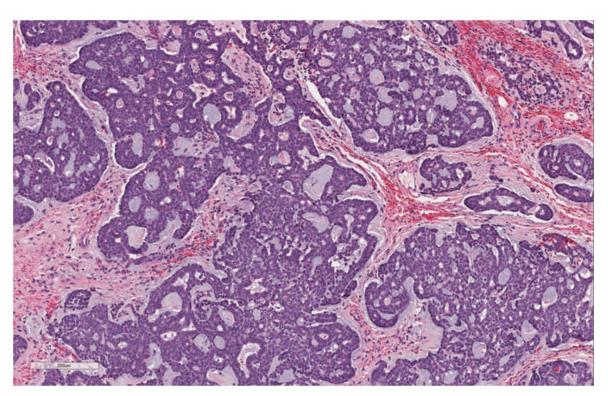


This tissue also forms glandular/tubular structures with lumen, and clearly represents the main lacrimal gland as well. However, instead of two well-ordered epi layers, the epi here is dense and chaotic. The multitude of small tubules is responsible for its 'Swiss cheese 'appearance.

What's the diagnosis?

Adenoid cystic carcinoma (ACC) is slightly more common in women. The median age at presentation is about 40 years. ACC has no pseudocapsule. Bone erosion is typical, and pain at presentation is the rule. Grows quickly vs slowly



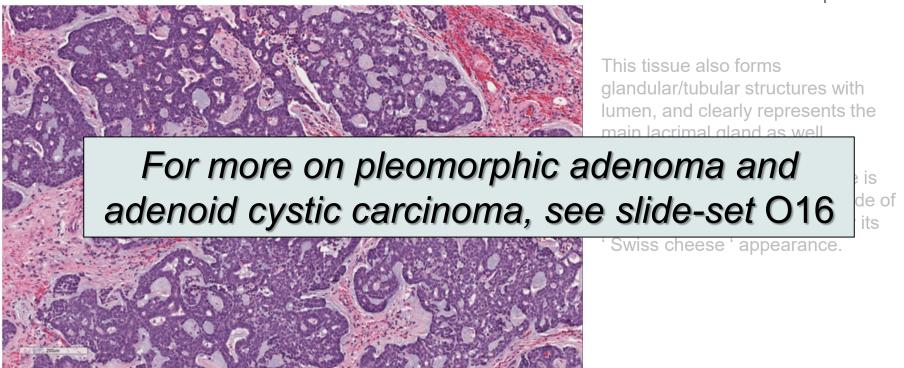


This tissue also forms glandular/tubular structures with lumen, and clearly represents the main lacrimal gland as well. However, instead of two well-ordered epi layers, the epi here is dense and chaotic. The multitude of small tubules is responsible for its 'Swiss cheese 'appearance.

What's the diagnosis?

Adenoid cystic carcinoma (ACC) is slightly more common in women. The median age at presentation is about 40 years. ACC has no pseudocapsule. Bone erosion is typical, and pain at presentation is the rule. Grows rapidly.

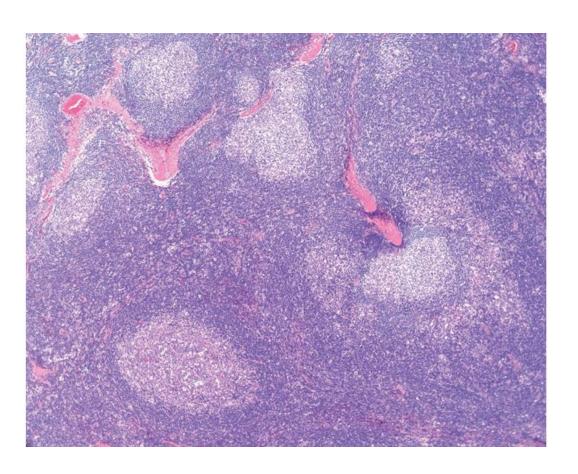




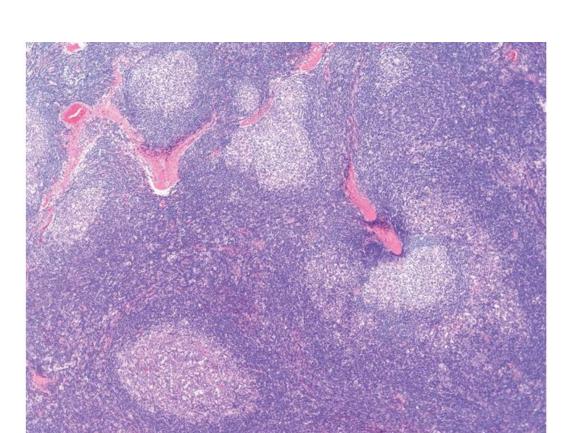
What's the diagnosis?

Adenoid cystic carcinoma (ACC) is slightly more common in women. The median age at presentation is about 40 years. ACC has no pseudocapsule. Bone erosion is typical, and pain at presentation is the rule. Grows rapidly.





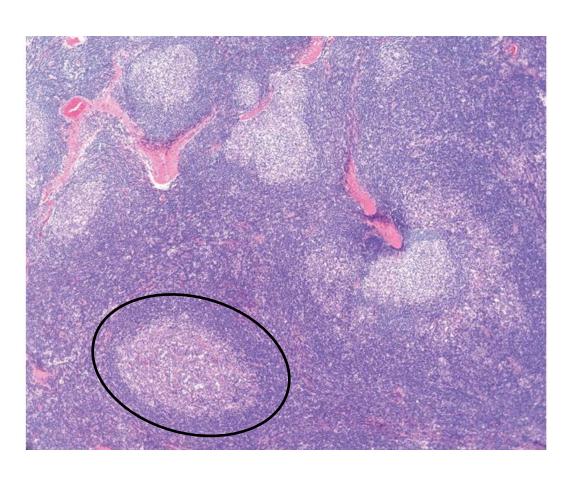
Hopefully, the appearance of this image screams very general to you.





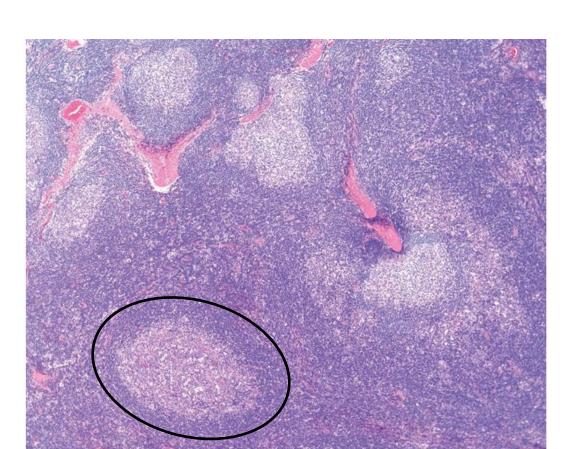
Hopefully, the appearance of this image screams *lymphoid* to you.





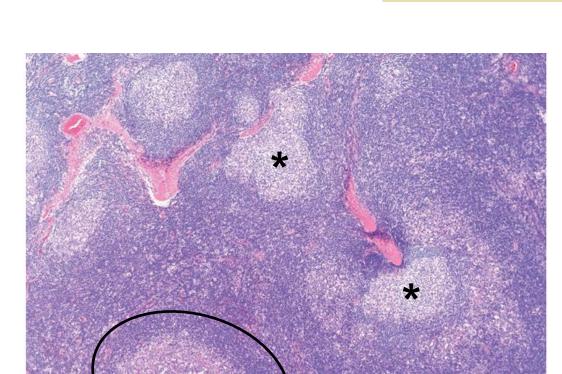
Hopefully, the appearance of this image screams *lymphoid* to you.

The tip-off is the presence of (here's one)





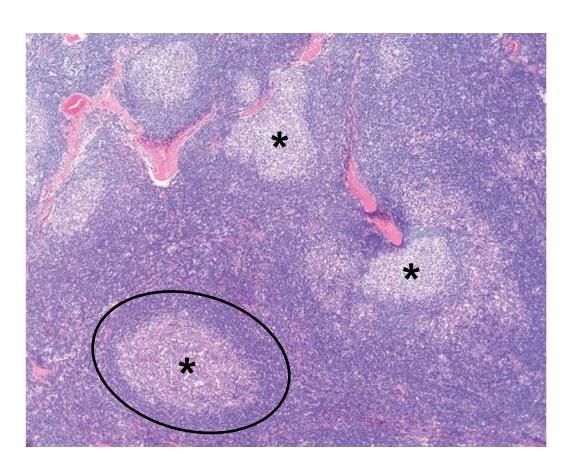
Hopefully, the appearance of this image screams *lymphoid* to you. The tip-off is the presence of follicles (here's one)





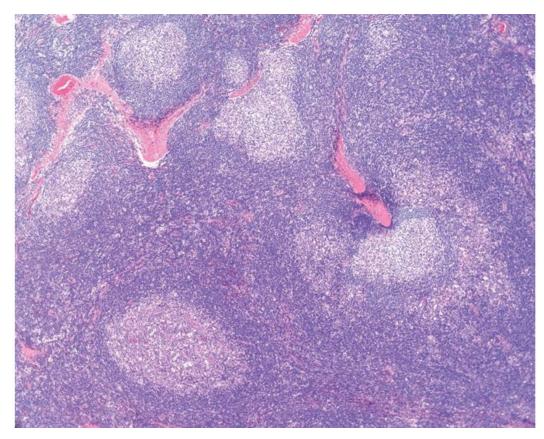
Hopefully, the appearance of this image screams *lymphoid* to you. The tip-off is the presence of follicles (here's one) with well-formed (asterisks).





Hopefully, the appearance of this image screams *lymphoid* to you. The tip-off is the presence of follicles (here's one) with well-formed germinal centers (asterisks).

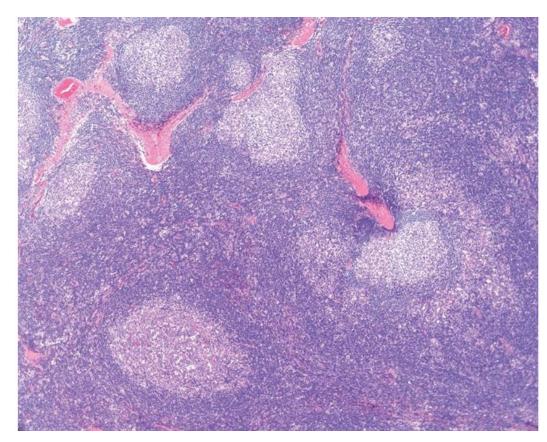




What's the diagnosis?

Hopefully, the appearance of this image screams *lymphoid* to you. The tip-off is the presence of follicles (here's one) with well-formed germinal centers (asterisks).



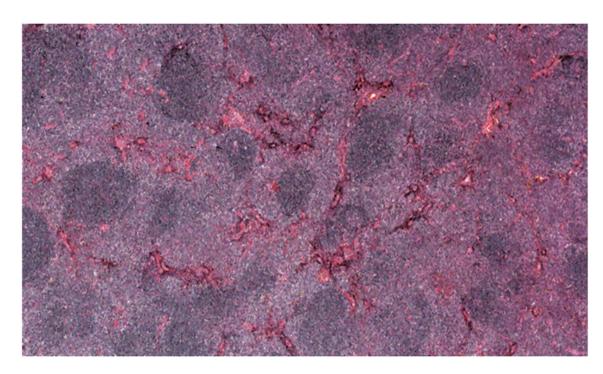


Hopefully, the appearance of this image screams *lymphoid* to you. The tip-off is the presence of follicles (here's one) with well-formed germinal centers (asterisks).

What's the diagnosis?

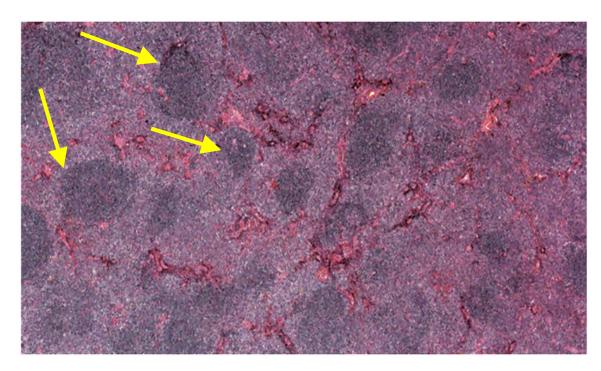
.





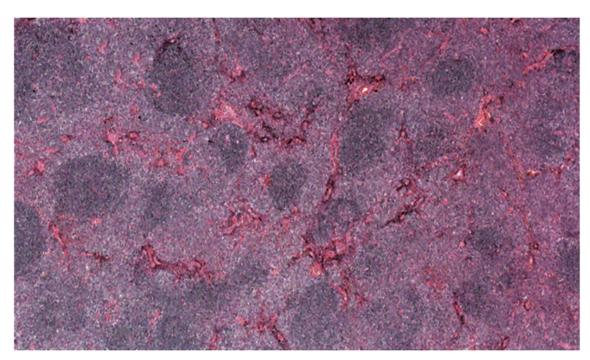
This one should also bring to mind the word *lymphoid*.





This one should also bring to mind the word *lymphoid*. However, note that its follicles (*arrows*) are much more haphazard-looking, and lack well-formed germinal centers.

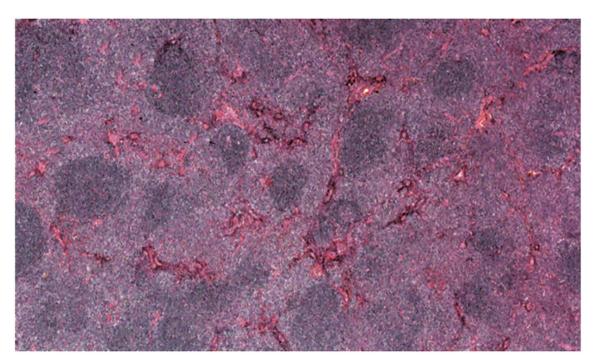




This one should also bring to mind the word *lymphoid*. However, note that its follicles (*arrows*) are much more haphazard-looking, and lack well-formed germinal centers.

What's the diagnosis?

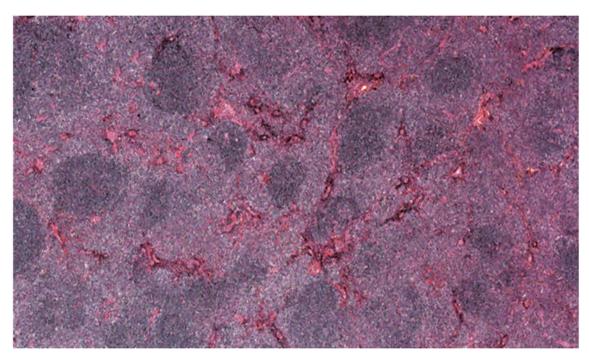




This one should also bring to mind the word *lymphoid*. However, note that its follicles (*arrows*) are much more haphazard-looking, and lack well-formed germinal centers.

What's the diagnosis?



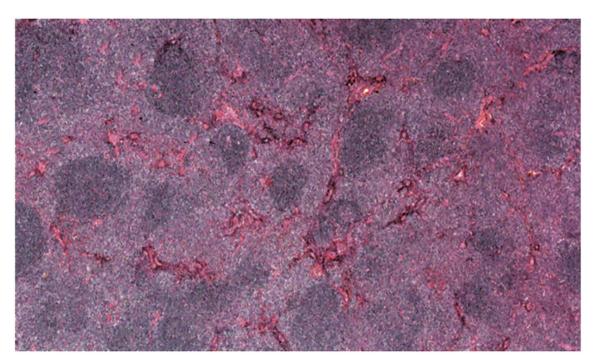


This one should also bring to mind the word *lymphoid*. However, note that its follicles (*arrows*) are much more haphazard-looking, and lack well-formed germinal centers.

What's the diagnosis?

Lymphoma. Most orbital lymphomas are non-Hogkins low-grade B-cell tumors.



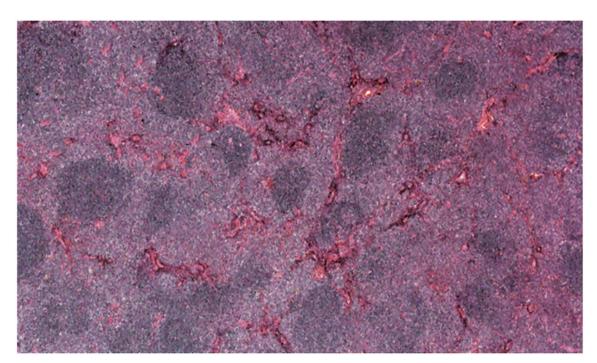


This one should also bring to mind the word *lymphoid*. However, note that its follicles (*arrows*) are much more haphazard-looking, and lack well-formed germinal centers.

What's the diagnosis?

Lymphoma. Most orbital lymphomas are non-Hogkins low-grade B-cell tumors. In general, their prognosis is good vs dire



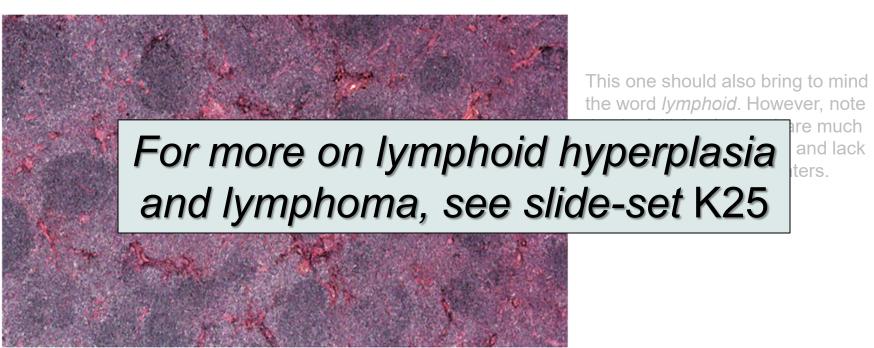


This one should also bring to mind the word *lymphoid*. However, note that its follicles (*arrows*) are much more haphazard-looking, and lack well-formed germinal centers.

What's the diagnosis?

Lymphoma. Most orbital lymphomas are non-Hogkins low-grade B-cell tumors. In general, their prognosis is good .

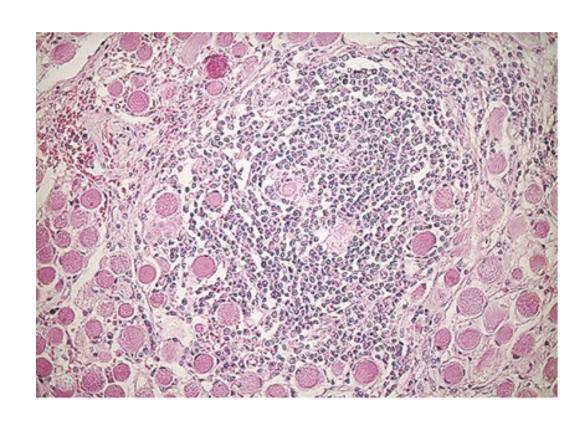




What's the diagnosis?

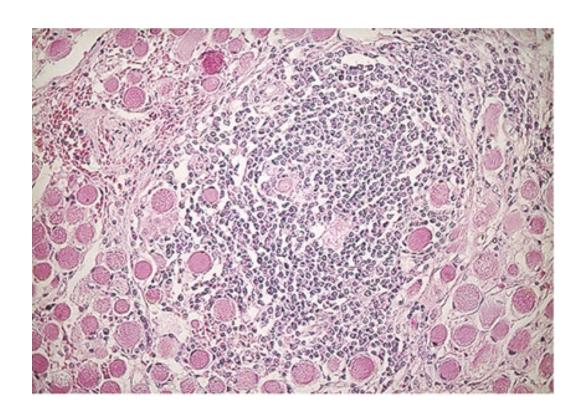
Lymphoma. Most orbital lymphomas are non-Hogkins low-grade B-cell tumors. In general, their prognosis is good .





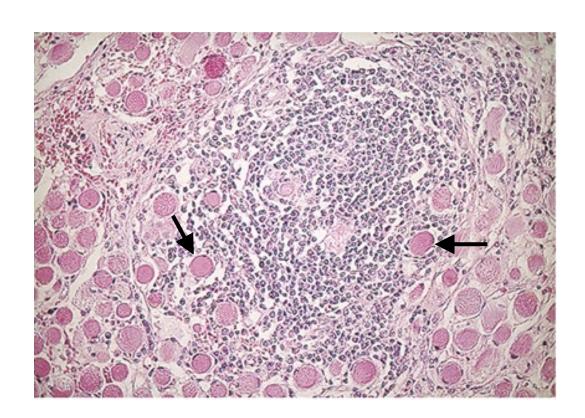
One of the first impressions you should take from this image is that there are a whole lotta cells present.





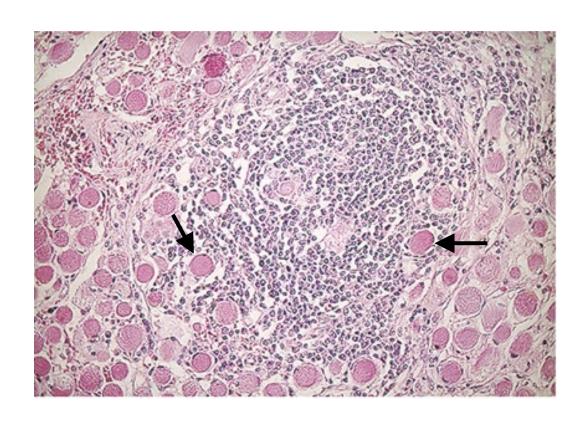
One of the first impressions you should take from this image is that there are a whole lotta inflammatory cells present.





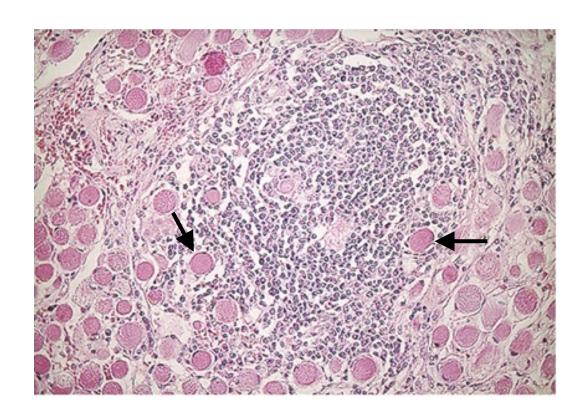
One of the first impressions you should take from this image is that there are a whole lotta inflammatory cells present. However, there's a repeated structure that's obviously not inflammatory—these things, which are two words





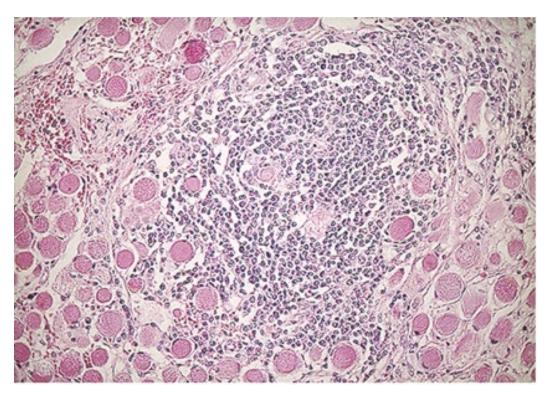
One of the first impressions you should take from this image is that there are a whole lotta inflammatory cells present. However, there's a repeated structure that's obviously not inflammatory—these things, which are muscle fibers.





One of the first impressions you should take from this image is that there are a whole lotta inflammatory cells present. However, there's a repeated structure that's obviously not inflammatory—these things, which are muscle fibers. As we're eyedocs, this means the image consists of inflammatory infiltrates within EOMs.

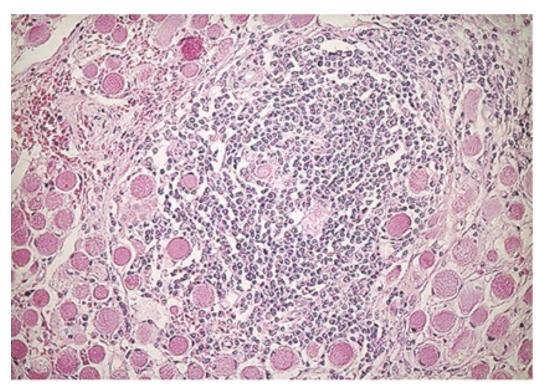




What's the diagnosis?

One of the first impressions you should take from this image is that there are a whole lotta inflammatory cells present. However, there's a repeated structure that's obviously not inflammatory—these things, which are muscle fibers. As we're eyedocs, this means the image consists of inflammatory infiltrates within EOMs.



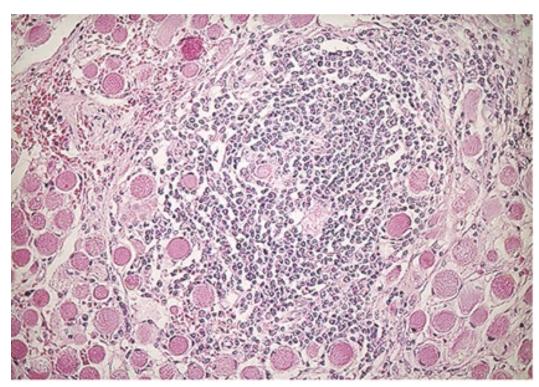


One of the first impressions you should take from this image is that there are a whole lotta inflammatory cells present. However, there's a repeated structure that's obviously not inflammatory—these things, which are muscle fibers. As we're eyedocs, this means the image consists of inflammatory infiltrates within EOMs.

(For the record, this is in fact NSOI)

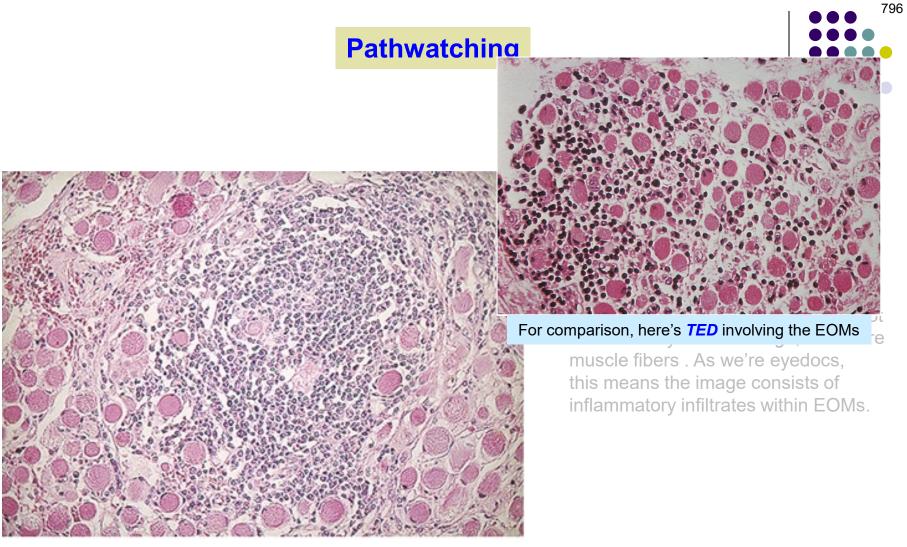
If you said nonspecific orbital inflammation (NSOI) or thyroid eye disease, give yourself a check.*



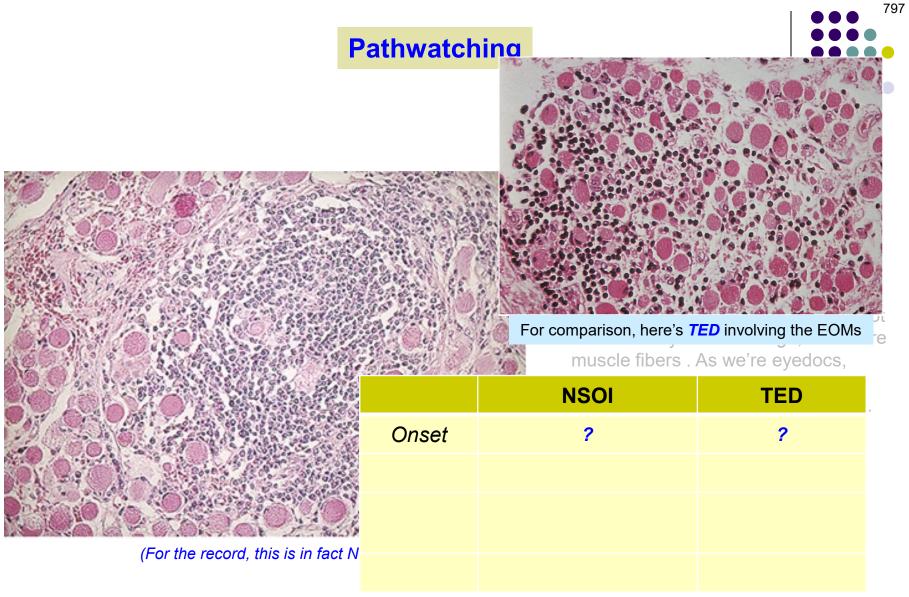


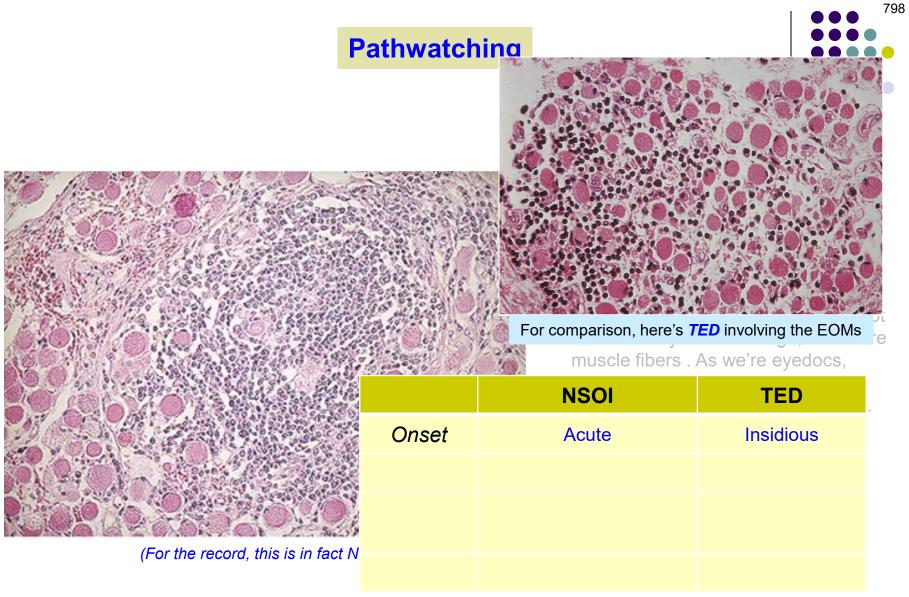
One of the first impressions you should take from this image is that there are a whole lotta inflammatory cells present. However, there's a repeated structure that's obviously not inflammatory—these things, which are muscle fibers. As we're eyedocs, this means the image consists of inflammatory infiltrates within EOMs.

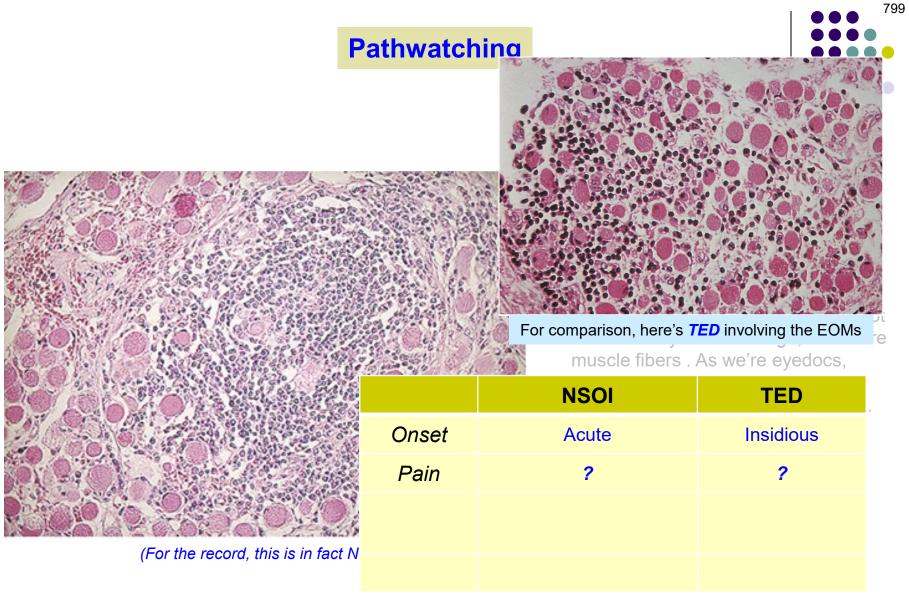
(For the record, this is in fact NSOI)

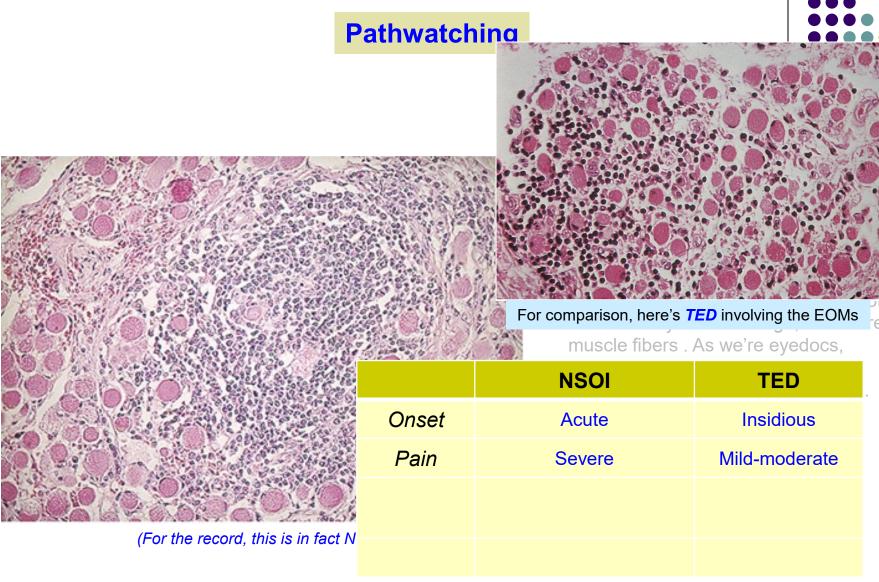


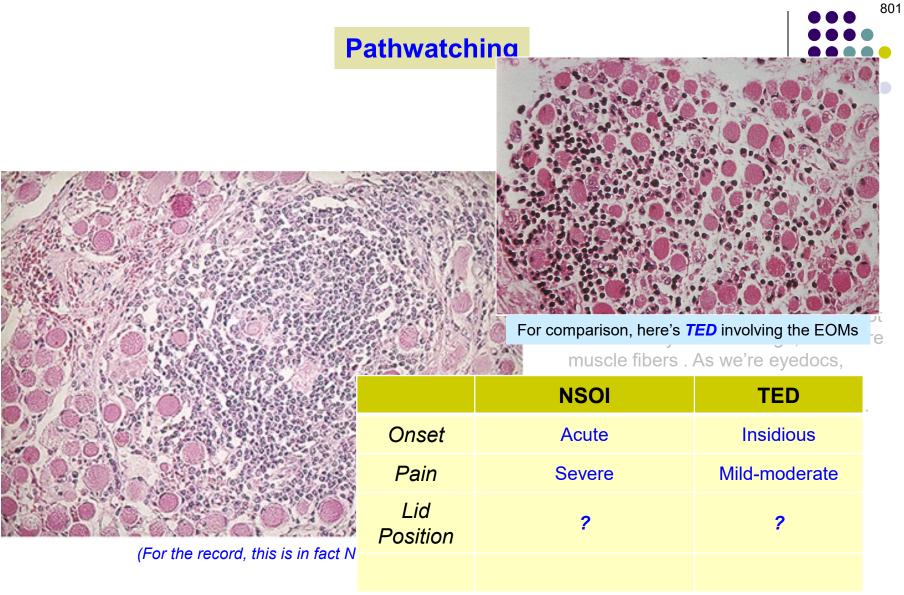
(For the record, this is in fact NSOI)

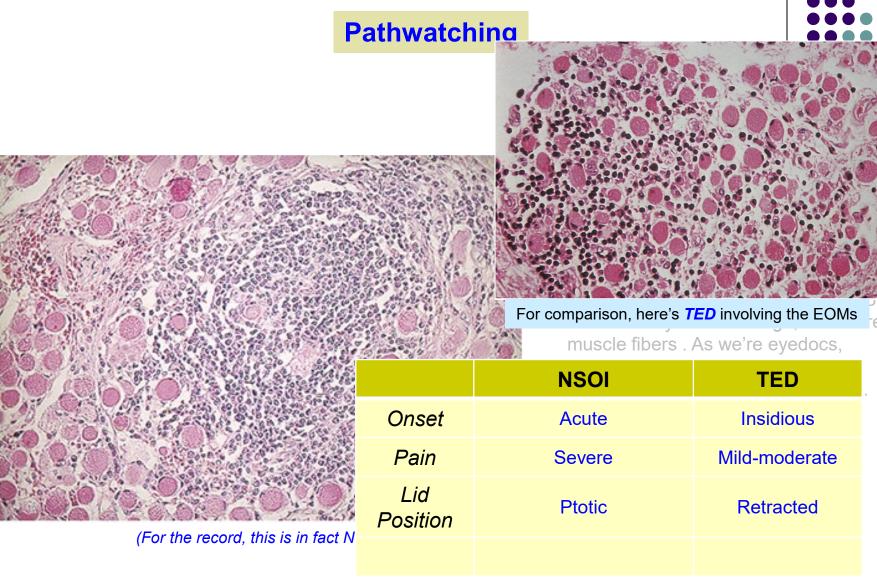


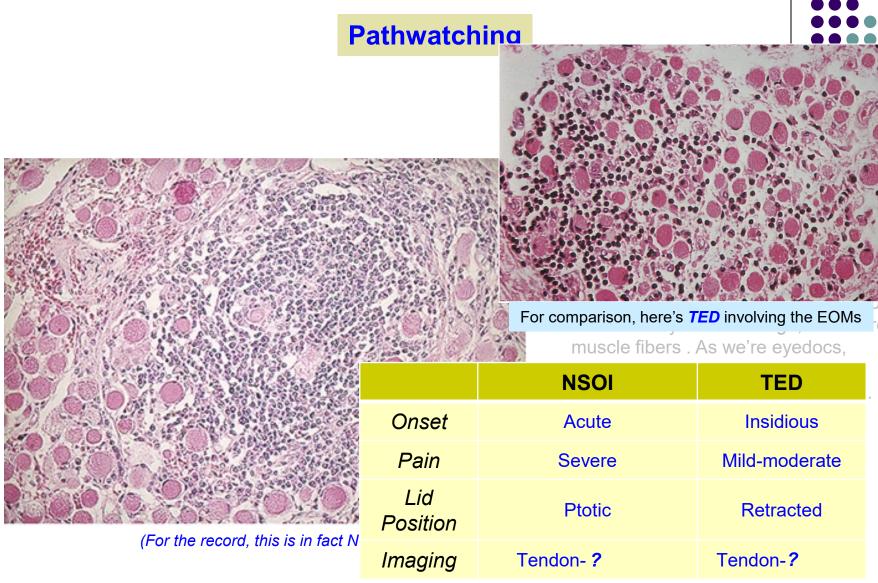


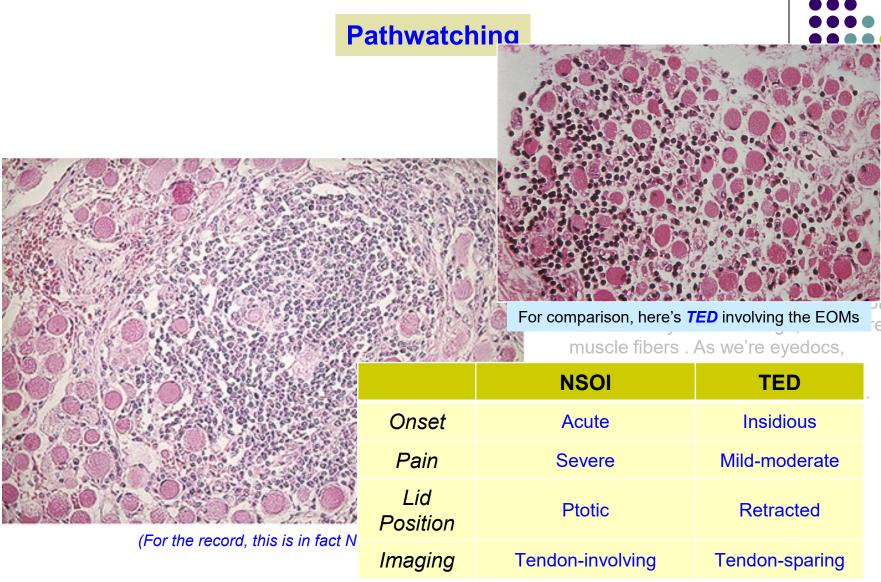


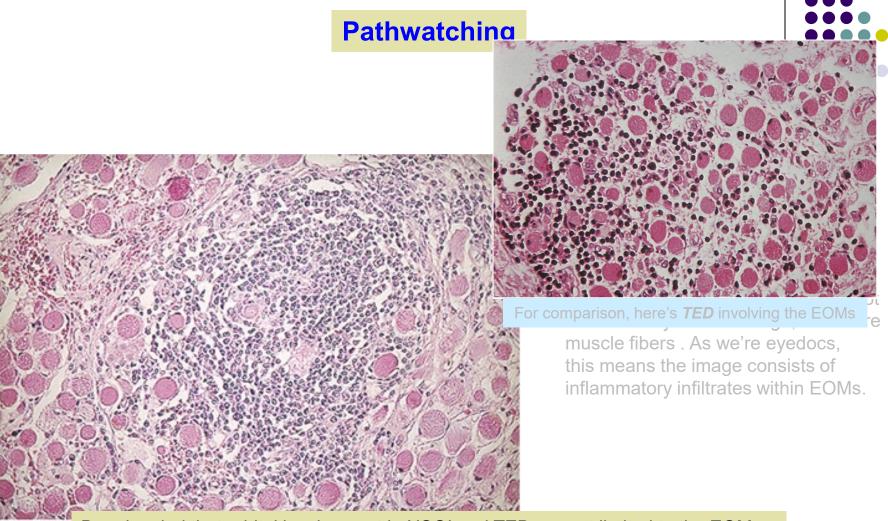






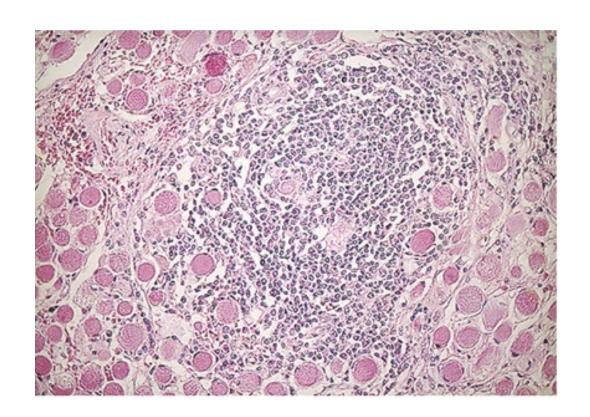






Bear in mind that orbital involvement in NSOI and TED are not limited to the EOM; any orbital structure can be affected (The point being, don't rely on the presence of muscle fibers on the slide to make these calls!)





One of the first impressions you should take from this image is that there are a whole lotta inflammatory cells present. However, there's a repeated structure that's obviously not inflammatory—these things, which are muscle fibers. As we're eyedocs, this means the image consists of inflammatory infiltrates within EOMs.

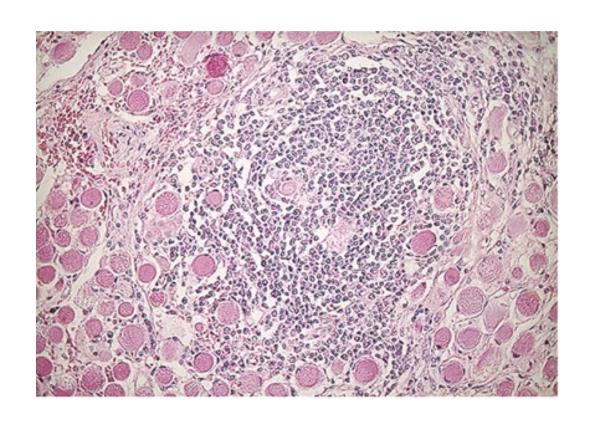
Finally, and circling back as promised: If you said

long name

Or two words

give yourself a check as well.

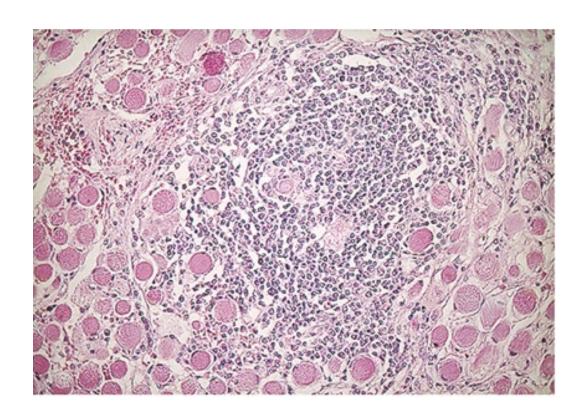




One of the first impressions you should take from this image is that there are a whole lotta inflammatory cells present. However, there's a repeated structure that's obviously not inflammatory—these things, which are muscle fibers. As we're eyedocs, this means the image consists of inflammatory infiltrates within EOMs.

Finally, and circling back as promised: If you said **IgG4-related orbital disease (IgG4-ROD)** or **Iymphoproliferative disease**, give yourself a check as well.

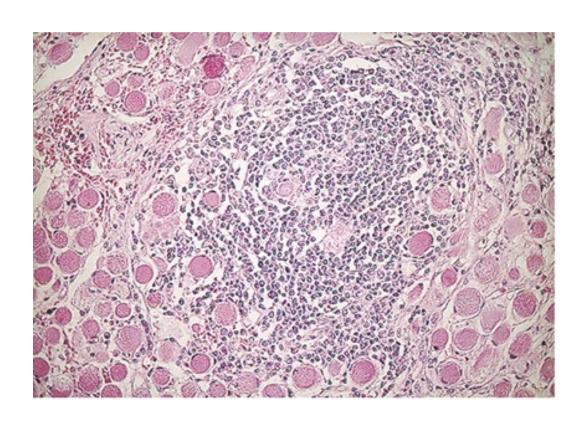




One of the first impressions you should take from this image is that there are a whole lotta inflammatory cells present. However, there's a repeated structure that's obviously not inflammatory—these things, which are muscle fibers. As we're eyedocs, this means the image consists of inflammatory infiltrates within EOMs.

Finally, and circling back as promised: If you said **IgG4-related orbital disease (IgG4-ROD)** or **Iymphoproliferative disease**, give yourself a check as well. (But a somewhat smaller one—while these conditions are mos def in the DDx, they are far more likely to involve the two words than the EOMs.)

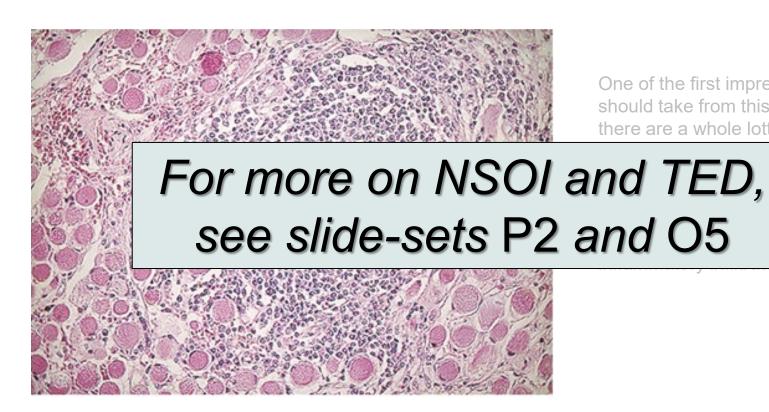




One of the first impressions you should take from this image is that there are a whole lotta inflammatory cells present. However, there's a repeated structure that's obviously not inflammatory—these things, which are muscle fibers. As we're eyedocs, this means the image consists of inflammatory infiltrates within EOMs.

Finally, and circling back as promised: If you said **IgG4-related orbital disease (IgG4-ROD)** or **Iymphoproliferative disease**, give yourself a check as well. (But a somewhat smaller one—while these conditions are mos def in the DDx, they are far more likely to involve the lacrimal gland than the EOMs.)



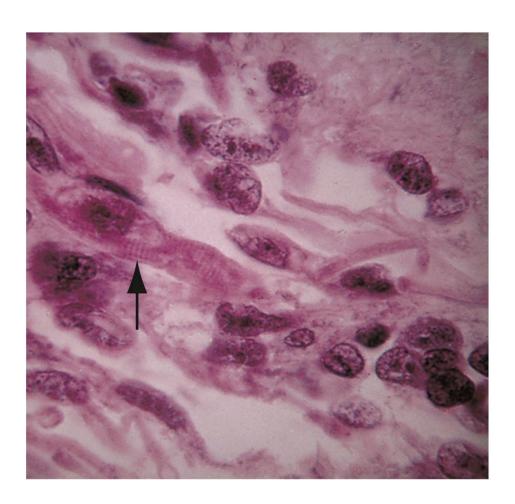


One of the first impressions you should take from this image is that there are a whole lotta inflammatory

> there's a s obviously not ings, which are evedocs, onsists of within EOMs.

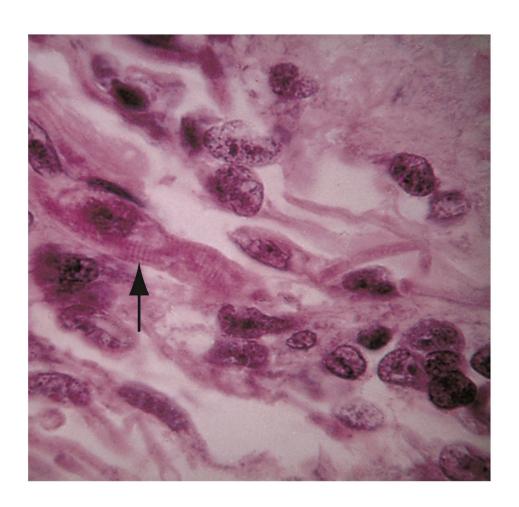
Finally, and circling back as promised: If you said **IgG4-related orbital disease (IgG4-ROD)** or lymphoproliferative disease, give yourself a check as well. (But a somewhat smaller one—while these conditions are mos def in the DDx, they are far more likely to involve the lacrimal gland than the EOMs.)





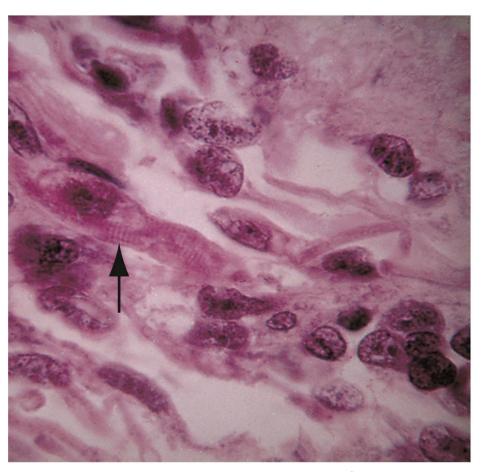
This field mark (*arrow*) is so difficult to see, I feel it would **have** to be pointed out on the OKAP. Look very carefully at it, then think way back to med school Path—what does it remind you of?





This field mark (arrow) is so difficult to see, I feel it would **have** to be pointed out on the OKAP. Look very carefully at it, then think way back to med school Path—what does it remind you of? If you said 'muscle fiber striations,' good on ya. Now look at the slide in general and note both its disordered appearance and the large, heavily stained nuclei. What does that make you think of?

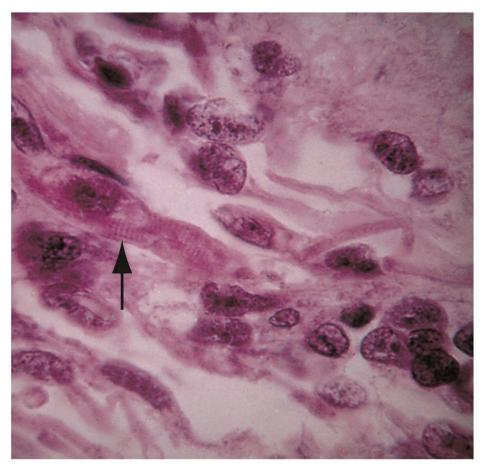




What's the diagnosis?

This field mark (arrow) is so difficult to see, I feel it would **have** to be pointed out on the OKAP. Look very carefully at it, then think way back to med school Path—what does it remind you of? If you said 'muscle fiber striations,' good on ya. Now look at the slide in general and note both its disordered appearance and the large, heavily stained nuclei. What does that make you think of? If you said malignancy, bingo. Now put your Ophtho hat back on—what sort of malignancy is this?





This field mark (arrow) is so difficult to see, I feel it would have to be pointed out on the OKAP. Look very carefully at it, then think way back to med school Path—what does it remind you of? If you said 'muscle fiber striations,' good on ya. Now look at the slide in general and note both its disordered appearance and the large, heavily stained nuclei. What does that make you think of? If you said malignancy, bingo. Now put your Ophtho hat back on—what sort of malignancy is this?

What's the diagnosis?

Rhabdomyosarcoma showing cross-striations (Z bands of actin-myosin complexes) within tumor cell cytoplasm



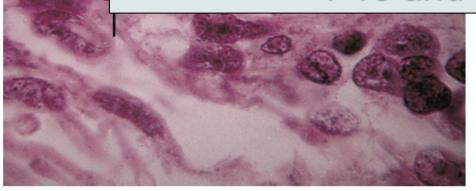


This field mark (*arrow*) is so difficult to see, I feel it would **have** to be pointed out on the OKAP. Look very carefully at it, then think way back to med school Path—what does it remind

striations,'
e in general
earance and
What does

For more on rhabdo, see slide-sets P15 and O13

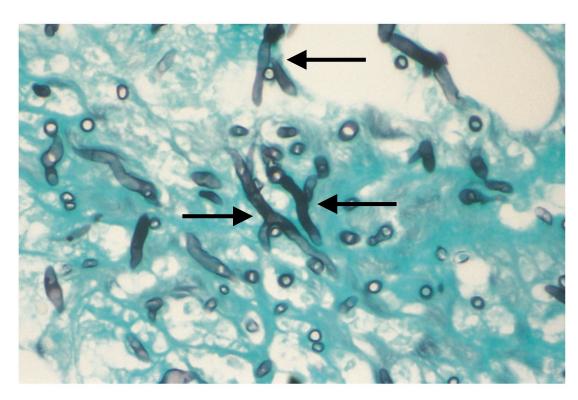
that make you think or: It you said malignancy, bingo. Now put your Ophtho hat back on—what sort of malignancy is this?



What's the diagnosis?

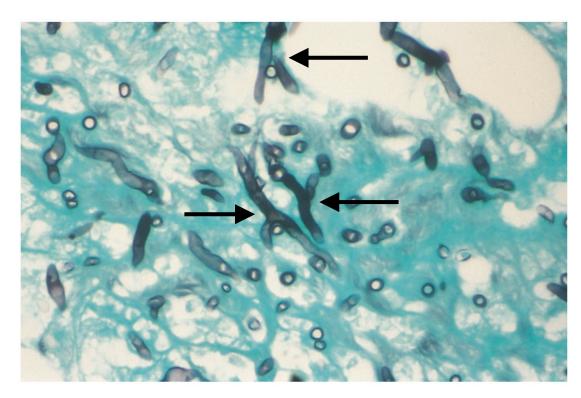
Rhabdomyosarcoma showing cross-striations (Z bands of actin-myosin complexes) within tumor cell cytoplasm





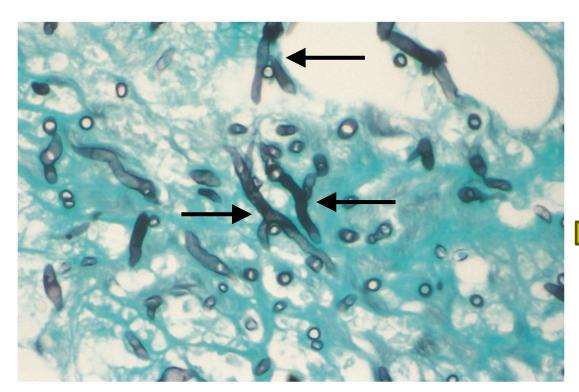
The field mark (*arrows*) for this condition is best remembered by describing it— 'they are, and they...'





The field mark (*arrows*) for this condition is best remembered by describing it— 'they are broad, and they branch.'

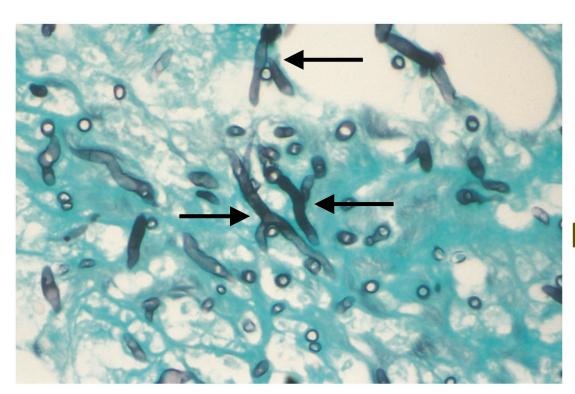




The field mark (*arrows*) for this condition is best remembered by describing it— 'they are broad, and they branch.'
There's a third word that goes with the first two; ie, when you think *broad* and *branch* the third should come immediately to mind. That word is

(Hint: It's an category dz).



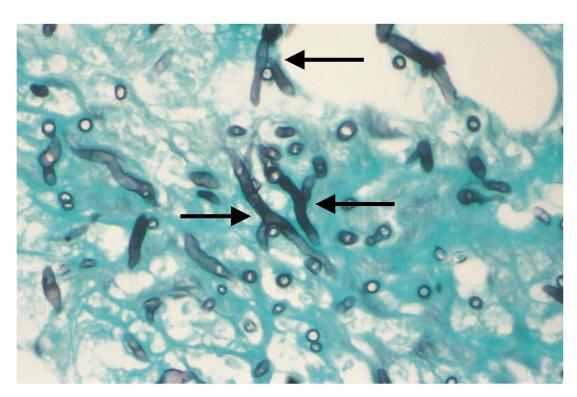


The field mark (*arrows*) for this condition is best remembered by describing it— 'they are broad, and they branch.'

There's a third word that goes with the first two; ie, when you think *broad* and *branch* the third should come immediately to mind. That word is

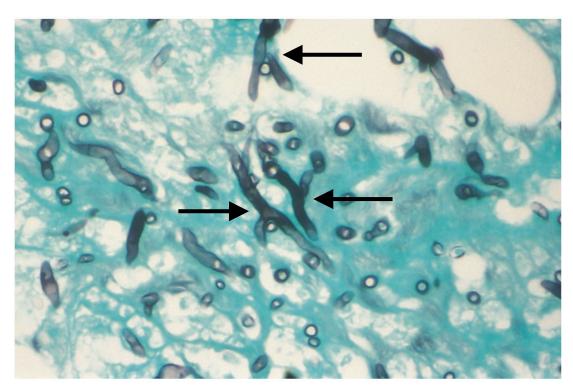
(Hint: It's an infectious dz).





The field mark (*arrows*) for this condition is best remembered by describing it— 'they are broad, and they branch.' There's a third word that goes with the first two; ie, when you think *broad* and *branch* the third should come immediately to mind. That word is *hyphae* (Hint: It's an infectious dz).

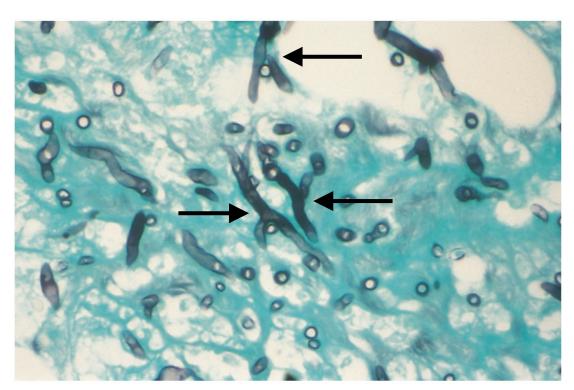




What's the diagnosis?

The field mark (*arrows*) for this condition is best remembered by describing it— 'they are broad, and they branch.' There's a third word that goes with the first two; ie, when you think *broad* and *branch* the third should come immediately to mind. That word is *hyphae* (Hint: It's an infectious dz).



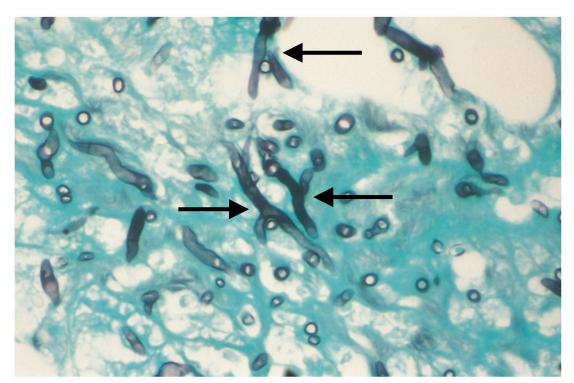


The field mark (*arrows*) for this condition is best remembered by describing it— 'they are broad, and they branch.' There's a third word that goes with the first two; ie, when you think *broad* and *branch* the third should come immediately to mind. That word is *hyphae* (Hint: It's an infectious dz).

What's the diagnosis?

Aspergillus infection showing broad branching fungal hyphae





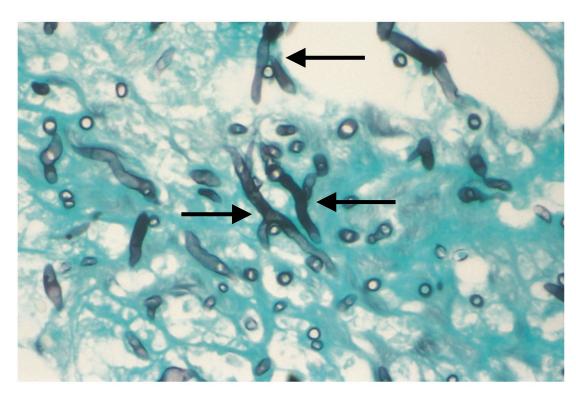
The field mark (*arrows*) for this condition is best remembered by describing it— 'they are broad, and they branch.' There's a third word that goes with the first two; ie, when you think *broad* and *branch* the third should come immediately to mind. That word is *hyphae* (Hint: It's an infectious dz).

What's the diagnosis?

Aspergillus infection showing broad branching fungal hyphae. Orbital aspergillosis comes in three forms:

- -- ? aspergillosis
- -- ? aspergillosis
- -- ? aspergillosis





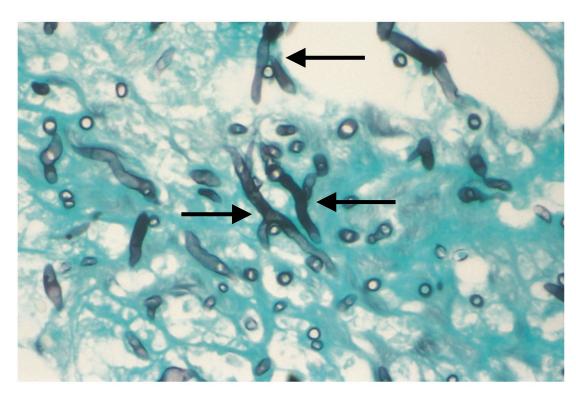
The field mark (*arrows*) for this condition is best remembered by describing it— 'they are broad, and they branch.' There's a third word that goes with the first two; ie, when you think *broad* and *branch* the third should come immediately to mind. That word is *hyphae* (Hint: It's an infectious dz).

What's the diagnosis?

Aspergillus infection showing broad branching fungal hyphae. Orbital aspergillosis comes in three forms:

- -- Invasive aspergillosis
- --Noninvasive aspergillosis
- -- Allergic aspergillosis





The field mark (arrows) for this condition is best remembered by describing it— 'they are broad, and they branch.' There's a third word that goes with the first two; ie, when you think broad and branch the third should come immediately to mind. That word is hyphae (Hint: It's an infectious dz).

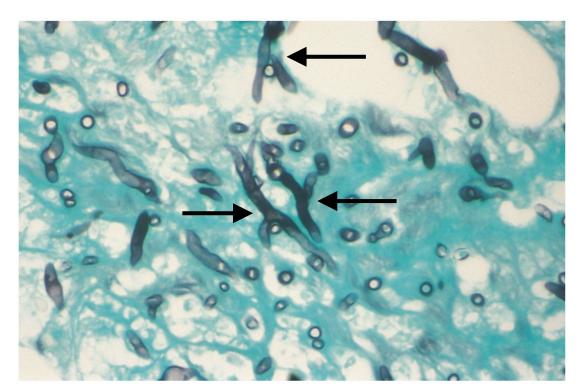
What's the diagnosis?

Aspergillus infection showing broad branching fungal hyphae. Orbital aspergillosis comes in three forms:

--Invasive aspergillosis: An broad category condition in general status pts

- --Noninvasive aspergillosis
- -- Allergic aspergillosis





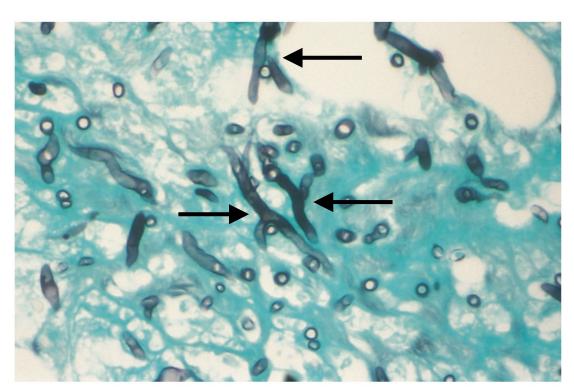
The field mark (arrows) for this condition is best remembered by describing it— 'they are broad, and they branch.' There's a third word that goes with the first two; ie, when you think broad and branch the third should come immediately to mind. That word is hyphae (Hint: It's an infectious dz).

What's the diagnosis?

Aspergillus infection showing broad branching fungal hyphae. Orbital aspergillosis comes in three forms: --Invasive aspergillosis: An infectious condition in immunocompromised pts

- --Noninvasive aspergillosis
- -- Allergic aspergillosis





The field mark (*arrows*) for this condition is best remembered by describing it— 'they are broad, and they branch.' There's a third word that goes with the first two; ie, when you think *broad* and *branch* the third should come immediately to mind. That word is *hyphae* (Hint: It's an infectious dz).

What's the diagnosis?

Aspergillus infection showing broad branching fungal hyphae. Orbital aspergillosis comes in three forms:

- --Invasive aspergillosis: An infectious condition in immunocompromised pts
- --Noninvasive aspergillosis: Characterized by the presence of an

one word

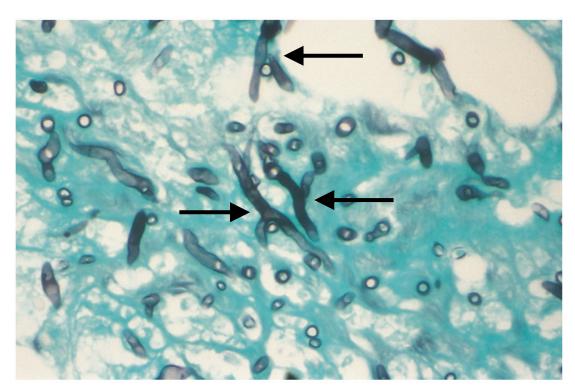
(aka a '

two words

....

-- Allergic aspergillosis





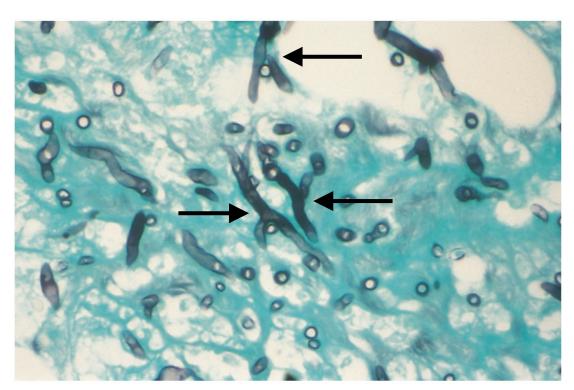
The field mark (*arrows*) for this condition is best remembered by describing it— 'they are broad, and they branch.' There's a third word that goes with the first two; ie, when you think *broad* and *branch* the third should come immediately to mind. That word is *hyphae* (Hint: It's an infectious dz).

What's the diagnosis?

Aspergillus infection showing broad branching fungal hyphae. Orbital aspergillosis comes in three forms:

- --Invasive aspergillosis: An infectious condition in immunocompromised pts
- --Noninvasive aspergillosis: Characterized by the presence of an aspergilloma (aka a ' fungal ball ')
- -- Allergic aspergillosis





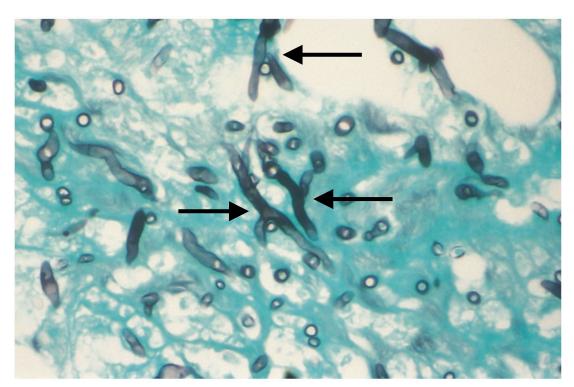
The field mark (*arrows*) for this condition is best remembered by describing it— 'they are broad, and they branch.' There's a third word that goes with the first two; ie, when you think *broad* and *branch* the third should come immediately to mind. That word is *hyphae* (Hint: It's an infectious dz).

What's the diagnosis?

Aspergillus infection showing broad branching fungal hyphae. Orbital aspergillosis comes in three forms:

- --Invasive aspergillosis: An infectious condition in immunocompromised pts
- --Noninvasive aspergillosis: Characterized by the presence of an aspergilloma (aka a ' fungal ball ')
- --Allergic aspergillosis: As named. Pts are usually





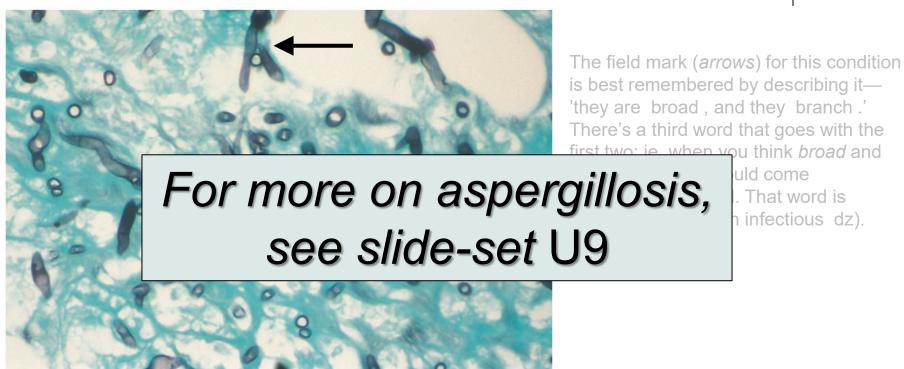
The field mark (*arrows*) for this condition is best remembered by describing it— 'they are broad, and they branch.' There's a third word that goes with the first two; ie, when you think *broad* and *branch* the third should come immediately to mind. That word is *hyphae* (Hint: It's an infectious dz).

What's the diagnosis?

Aspergillus infection showing broad branching fungal hyphae. Orbital aspergillosis comes in three forms:

- --Invasive aspergillosis: An infectious condition in immunocompromised pts
- --Noninvasive aspergillosis: Characterized by the presence of an aspergilloma (aka a ' fungal ball ')
- --Allergic aspergillosis: As named. Pts are usually atopic



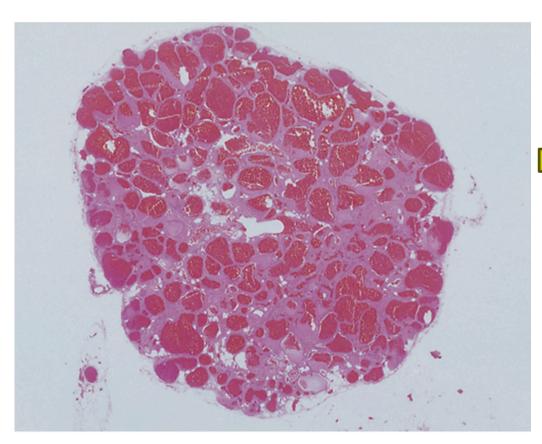


What's the diagnosis?

Aspergillus infection showing broad branching fungal hyphae. Orbital aspergillosis comes in three forms:

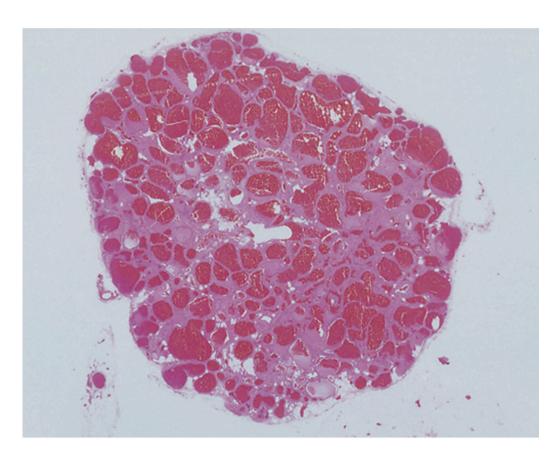
- --Invasive aspergillosis: An infectious condition in immunocompromised pts
- --Noninvasive aspergillosis: Characterized by the presence of an aspergilloma (aka a 'fungal ball ')
- --Allergic aspergillosis: As named. Pts are usually atopic





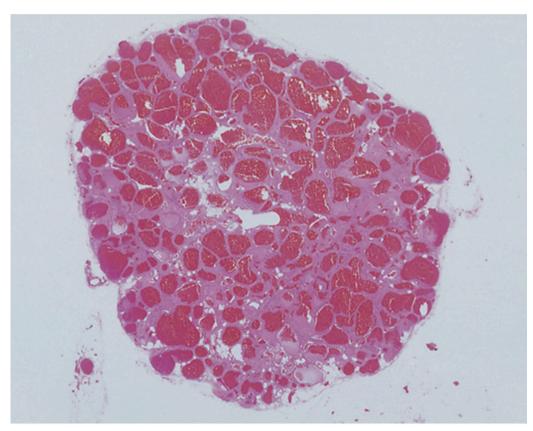
The striking thing about this mass is that it is comprised chiefly of numerous large two-words spaces. That's all you need to know to ID it.





The striking thing about this mass is that it is comprised chiefly of numerous large blood-filled spaces. That's all you need to know to ID it.

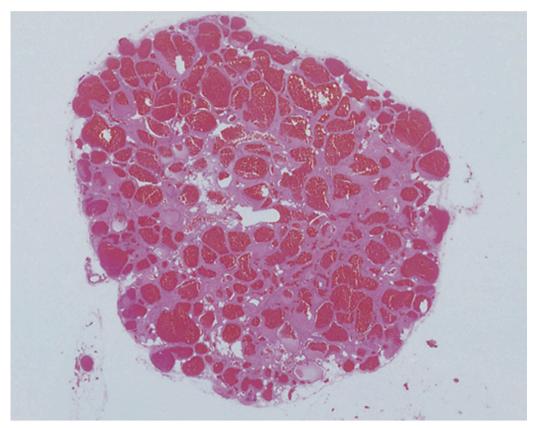




What's the diagnosis?

The striking thing about this mass is that it is comprised chiefly of numerous large blood-filled spaces. That's all you need to know to ID it.





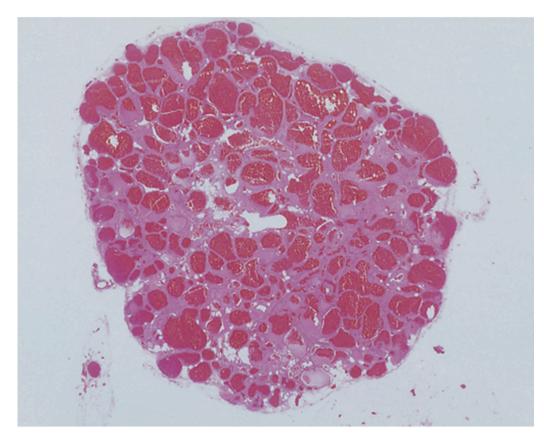
The striking thing about this mass is that it is comprised chiefly of numerous large blood-filled spaces. That's all you need to know to ID it.

What's the diagnosis?

Cavernous hemangioma (aka

three words



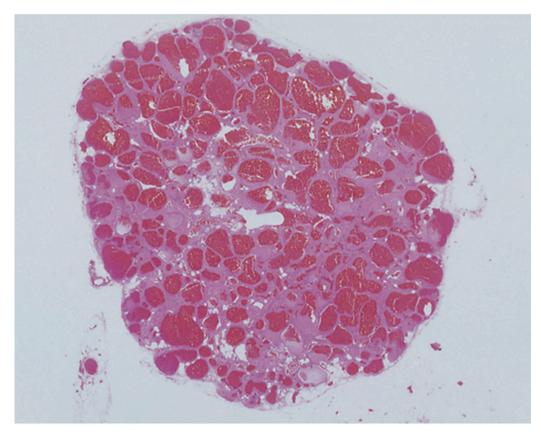


The striking thing about this mass is that it is comprised chiefly of numerous large blood-filled spaces. That's all you need to know to ID it.

What's the diagnosis?

Cavernous hemangioma (aka cavernous venous malformation)





The striking thing about this mass is that it is comprised chiefly of numerous large blood-filled spaces. That's all you need to know to ID it.

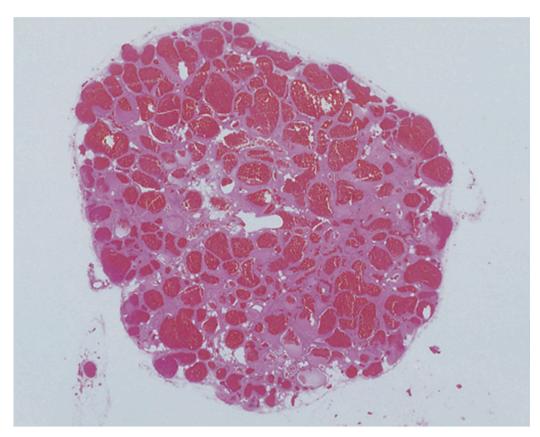
What's the diagnosis?

Cavernous hemangioma (aka cavernous venous malformation) is the least primary orbital lesion in adults



common



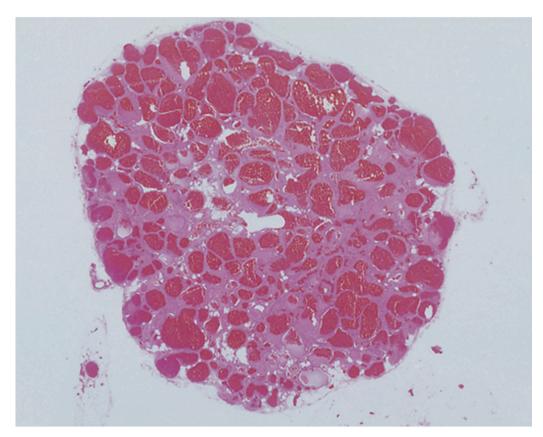


The striking thing about this mass is that it is comprised chiefly of numerous large blood-filled spaces. That's all you need to know to ID it.

What's the diagnosis?

Cavernous hemangioma (aka cavernous venous malformation) is the most common primary orbital lesion in adults



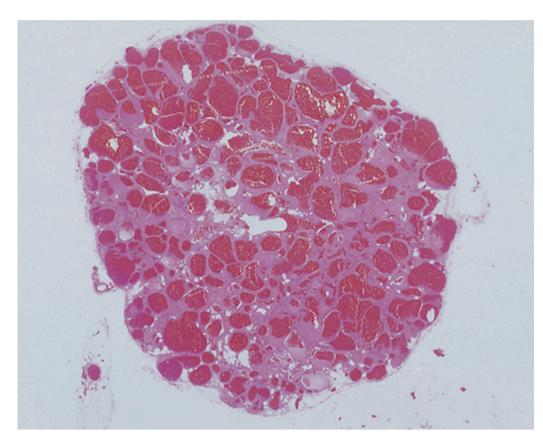


The striking thing about this mass is that it is comprised chiefly of numerous large blood-filled spaces. That's all you need to know to ID it.

What's the diagnosis?

Cavernous hemangioma (aka cavernous venous malformation) is the most common primary orbital lesion in adults (usually the range decade).



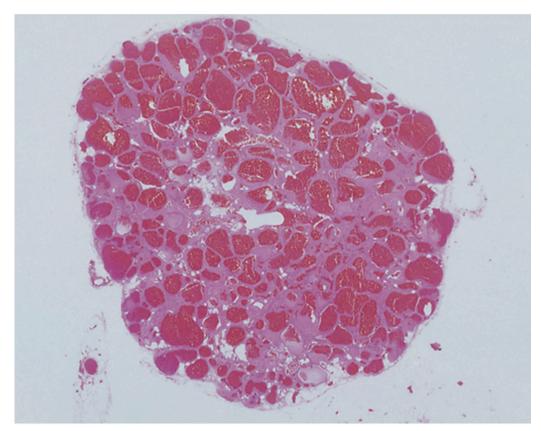


The striking thing about this mass is that it is comprised chiefly of numerous large blood-filled spaces. That's all you need to know to ID it.

What's the diagnosis?

Cavernous hemangioma (aka *cavernous venous malformation*) is the most common primary orbital lesion in adults (usually the 4th to 5th decade).



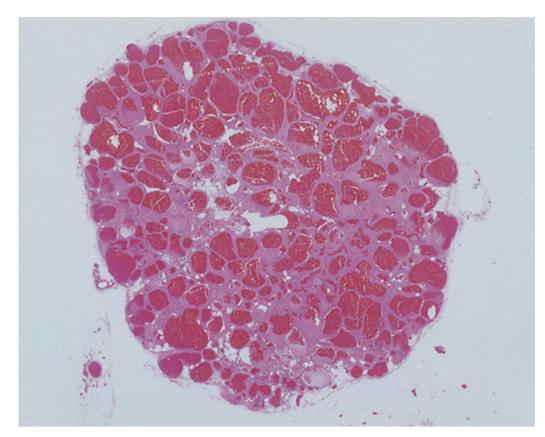


The striking thing about this mass is that it is comprised chiefly of numerous large blood-filled spaces. That's all you need to know to ID it.

What's the diagnosis?

Cavernous hemangioma (aka cavernous venous malformation) is the most common primary orbital lesion in adults (usually the 4th to 5th decade). More common in men vs women



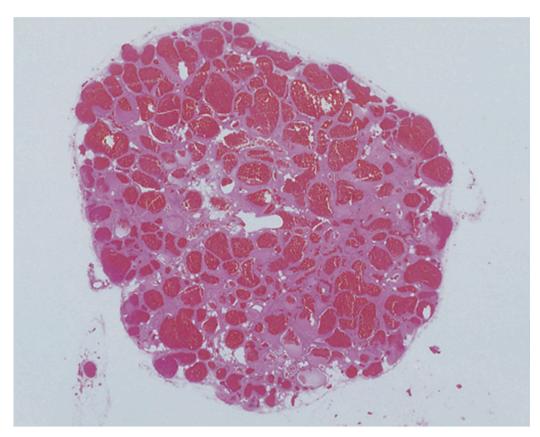


The striking thing about this mass is that it is comprised chiefly of numerous large blood-filled spaces. That's all you need to know to ID it.

What's the diagnosis?

Cavernous hemangioma (aka *cavernous venous malformation*) is the most common primary orbital lesion in adults (usually the 4th to 5th decade). More common in women .



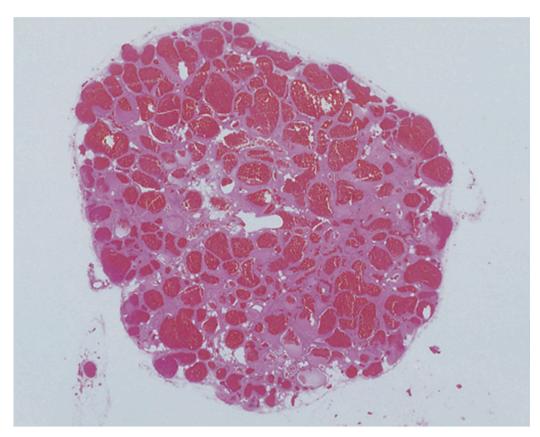


The striking thing about this mass is that it is comprised chiefly of numerous large blood-filled spaces. That's all you need to know to ID it.

What's the diagnosis?

Cavernous hemangioma (aka cavernous venous malformation) is the most common primary orbital lesion in adults (usually the 4th to 5th decade). More common in women. Presents with slowly vs progressive and painful vs proptosis.



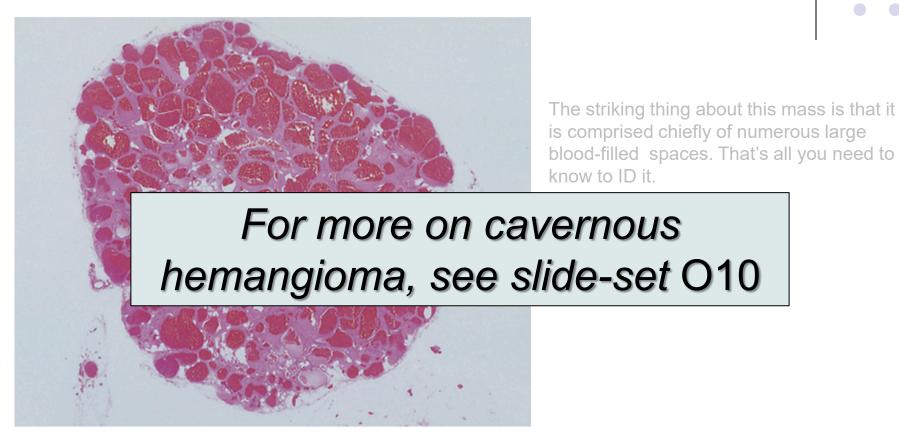


The striking thing about this mass is that it is comprised chiefly of numerous large blood-filled spaces. That's all you need to know to ID it.

What's the diagnosis?

Cavernous hemangioma (aka *cavernous venous malformation*) is the most common primary orbital lesion in adults (usually the 4th to 5th decade). More common in women. Presents with slowly progressive and painless proptosis.

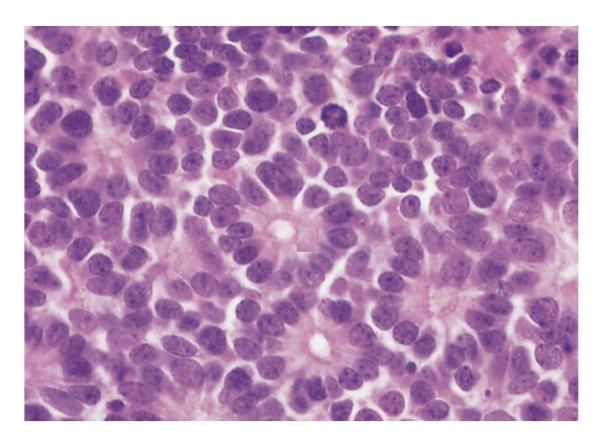




What's the diagnosis?

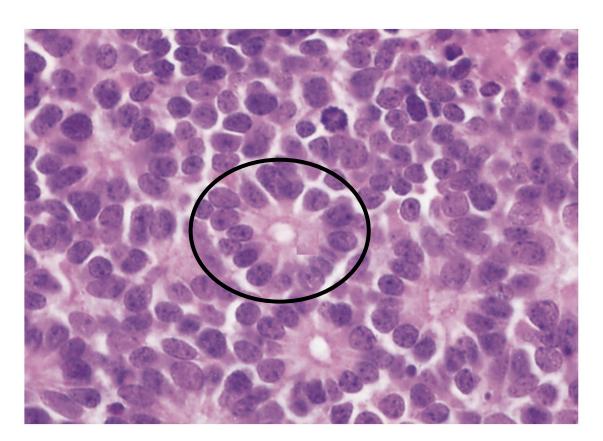
Cavernous hemangioma (aka *cavernous venous malformation*) is the most common primary orbital lesion in adults (usually the 4^{th} to 5^{th} decade). More common in women . Presents with slowly progressive and painless proptosis.





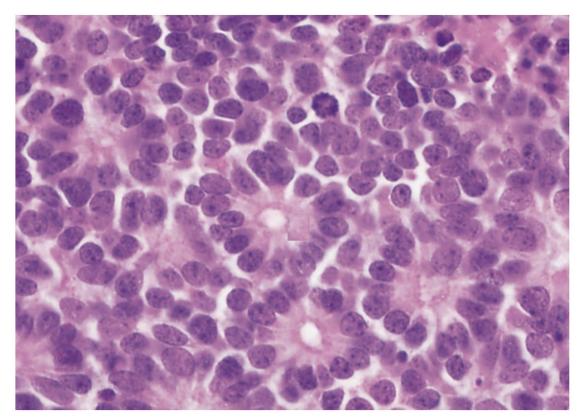
There is a classic finding here of a high-profile ophthalmic condition—can you find it?





There is a classic finding here of a high-profile ophthalmic condition—can you find it? **This** is it.

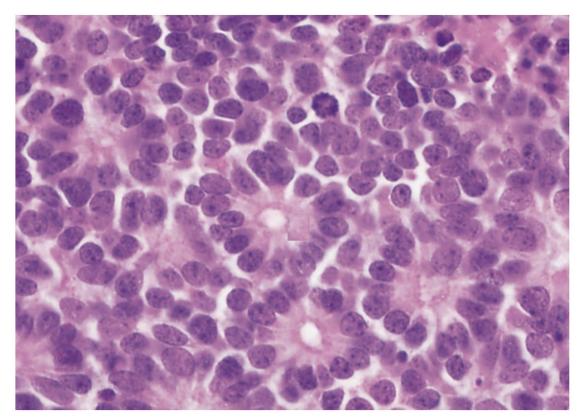




There is a classic finding here of a high-profile ophthalmic condition—can you find it? **This** is it.

What's the diagnosis?





There is a classic finding here of a high-profile ophthalmic condition—can you find it? **This** is it.

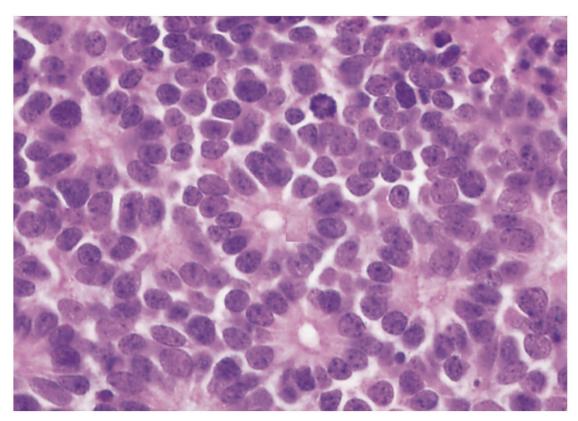
What's the diagnosis?

two words

rosettes in

disease



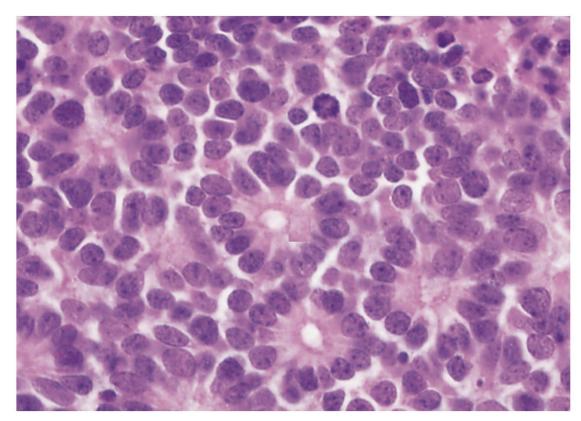


There is a classic finding here of a high-profile ophthalmic condition—can you find it? **This** is it.

What's the diagnosis?

Flexner-Wintersteiner rosettes in retinoblastoma (Rb) .





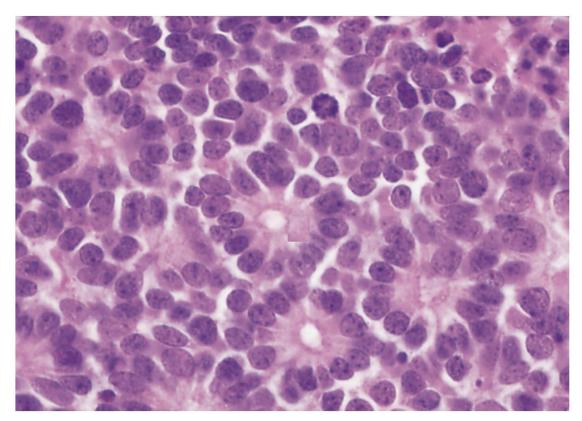
There is a classic finding here of a high-profile ophthalmic condition—can you find it? **This** is it.

What's the diagnosis?

Flexner-Wintersteiner rosettes in retinoblastoma (Rb) Is vs Is not





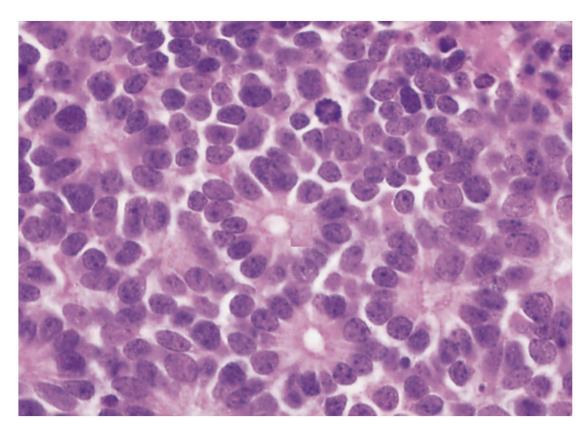


There is a classic finding here of a high-profile ophthalmic condition—can you find it? **This** is it.

What's the diagnosis?

Flexner-Wintersteiner rosettes in retinoblastoma (Rb) . Not pathognomonic for Rb



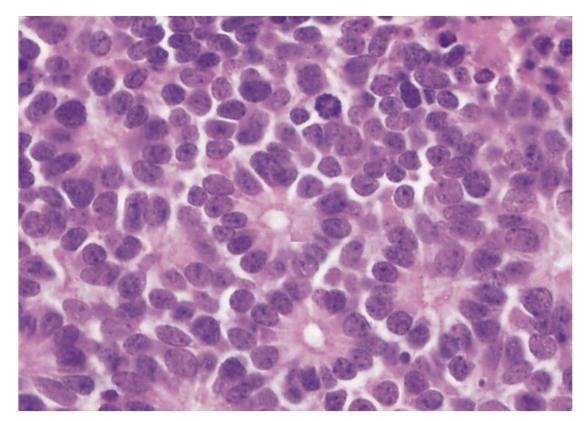


There is a classic finding here of a high-profile ophthalmic condition—can you find it? **This** is it.

What's the diagnosis?

Flexner-Wintersteiner rosettes in retinoblastoma (Rb). Not pathognomonic for Rb, but they are only rarely found in other general cell type tumors.



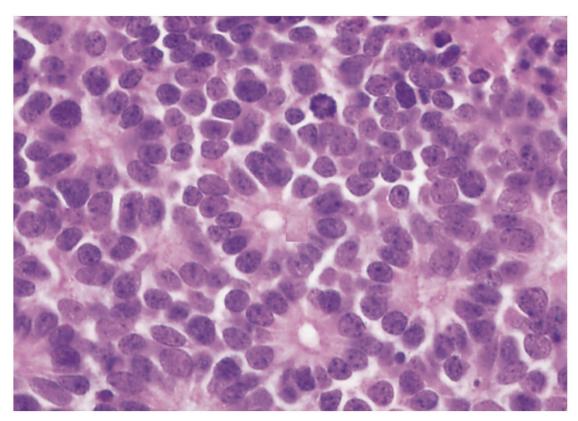


There is a classic finding here of a high-profile ophthalmic condition—can you find it? **This** is it.

What's the diagnosis?

Flexner-Wintersteiner rosettes in **retinoblastoma** (Rb). Not pathognomonic for Rb, but they are only rarely found in other neuroblastic tumors.



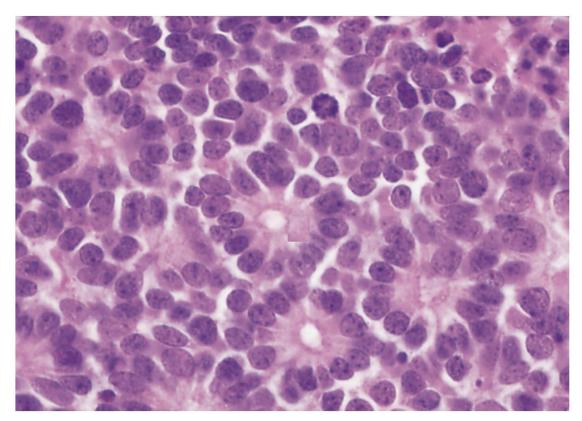


There is a classic finding here of a high-profile ophthalmic condition—can you find it? **This** is it.

What's the diagnosis?

Flexner-Wintersteiner rosettes in retinoblastoma (Rb). Not pathognomonic for Rb, but they are only rarely found in other neuroblastic tumors. Represents of the tumor, ie, an attempt to form mature retinal tissue.



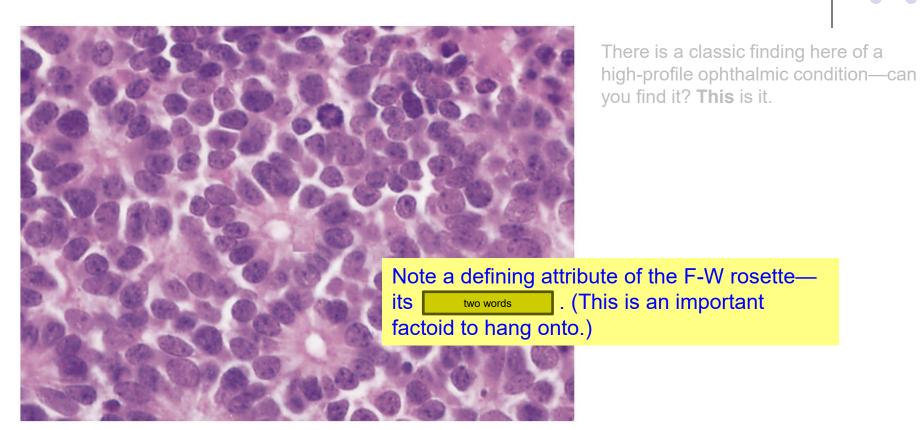


There is a classic finding here of a high-profile ophthalmic condition—can you find it? **This** is it.

What's the diagnosis?

Flexner-Wintersteiner rosettes in **retinoblastoma** (Rb). Not pathognomonic for Rb, but they are only rarely found in other neuroblastic tumors. Represents differentiation of the tumor, ie, an attempt to form mature retinal tissue.

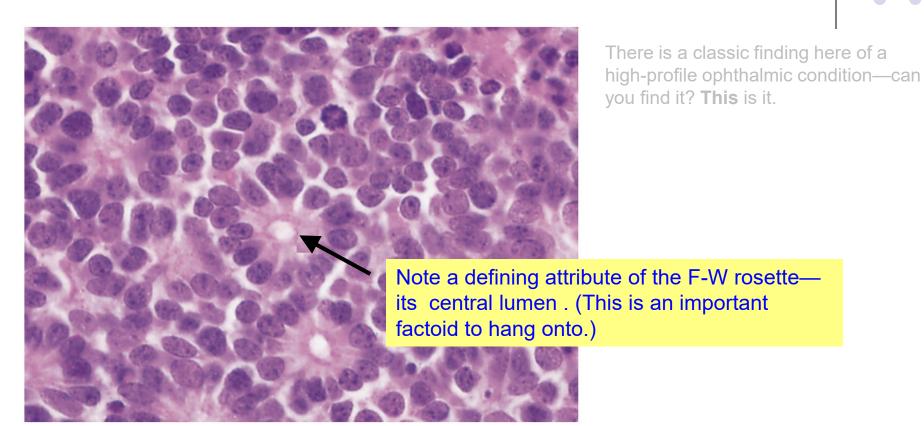




What's the diagnosis?

Flexner-Wintersteiner rosettes in **retinoblastoma** (Rb). Not pathognomonic for Rb, but they are only rarely found in other neuroblastic tumors. Represents differentiation of the tumor, ie, an attempt to form mature retinal tissue.

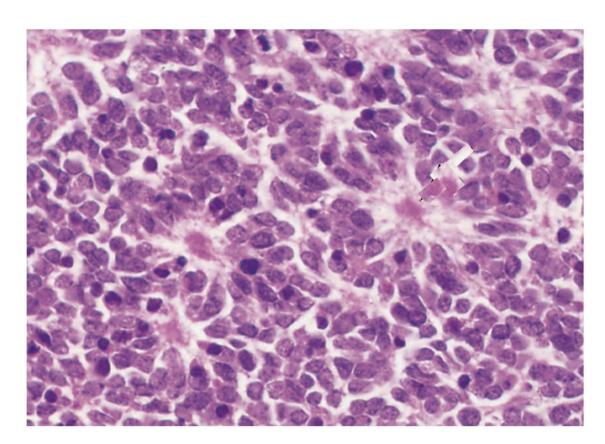




What's the diagnosis?

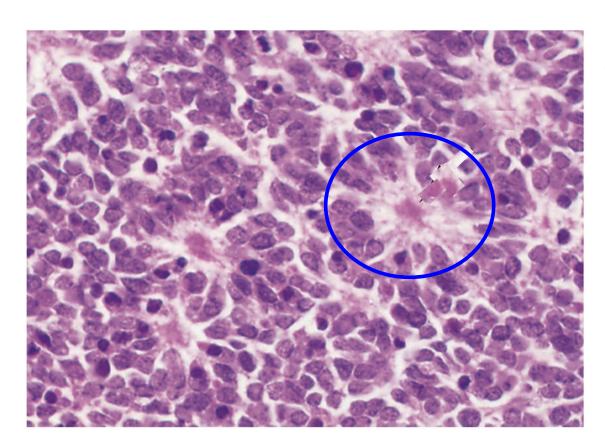
Flexner-Wintersteiner rosettes in **retinoblastoma** (Rb). Not pathognomonic for Rb, but they are only rarely found in other neuroblastic tumors. Represents differentiation of the tumor, ie, an attempt to form mature retinal tissue.





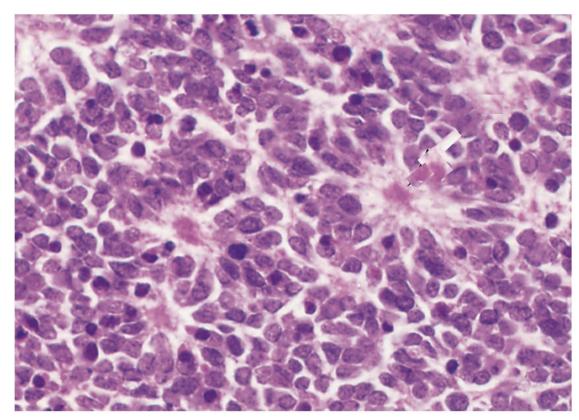
Another classic Rb finding (albeit neither as infamous nor as easy to spot as F-W rosettes) is depicted here—can you spot it?





Another classic Rb finding (albeit neither as infamous nor as easy to spot as F-W rosettes) is depicted here—can you spot it? Here it is.

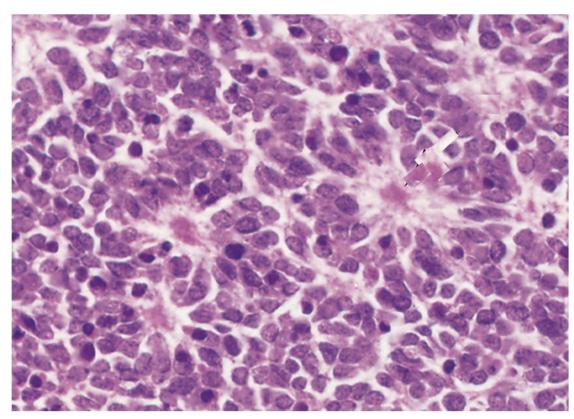




Another classic Rb finding (albeit neither as infamous nor as easy to spot as F-W rosettes) is depicted here—can you spot it? Here it is.

What's the finding?





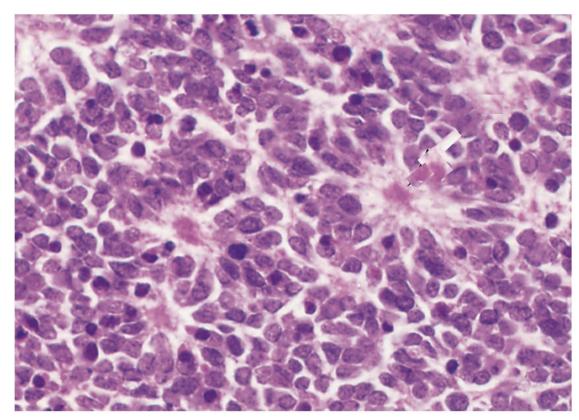
Another classic Rb finding (albeit neither as infamous nor as easy to spot as F-W rosettes) is depicted here—can you spot it? **Here** it is.

What's the finding?

two words

rosettes in retinoblastoma (Rb).



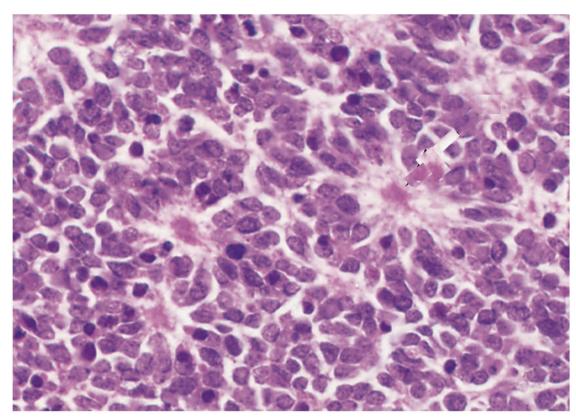


Another classic Rb finding (albeit neither as infamous nor as easy to spot as F-W rosettes) is depicted here—can you spot it? **Here** it is.

What's the finding?

Homer Wright rosettes in retinoblastoma (Rb).

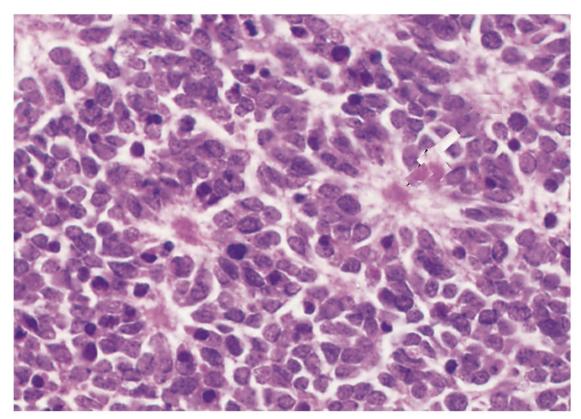




Another classic Rb finding (albeit neither as infamous nor as easy to spot as F-W rosettes) is depicted here—can you spot it? **Here** it is.

What's the finding?



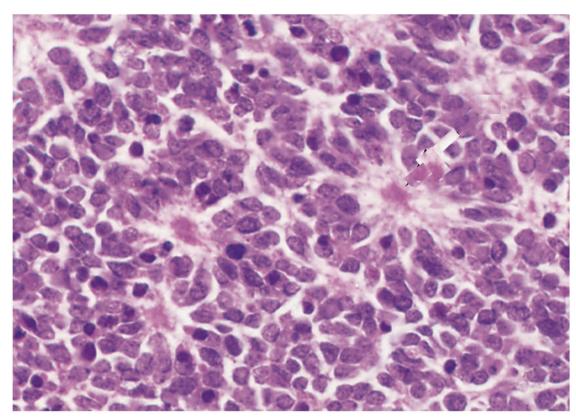


Another classic Rb finding (albeit neither as infamous nor as easy to spot as F-W rosettes) is depicted here—can you spot it? **Here** it is.

What's the finding?

Homer Wright rosettes in retinoblastoma (Rb). Not pathognomonic for Rb



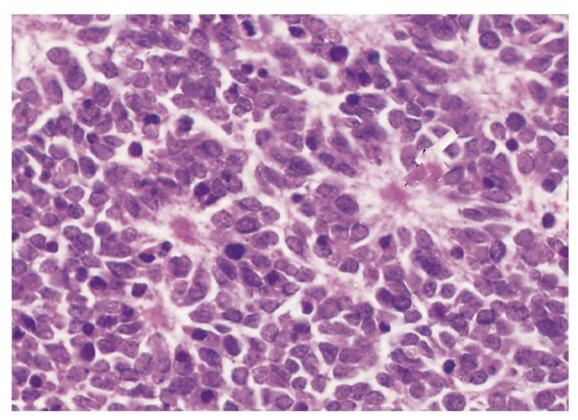


Another classic Rb finding (albeit neither as infamous nor as easy to spot as F-W rosettes) is depicted here—can you spot it? **Here** it is.

What's the finding?

Homer Wright rosettes in retinoblastoma (Rb). Not pathognomonic for Rb; they are found in other general cell type tumors.



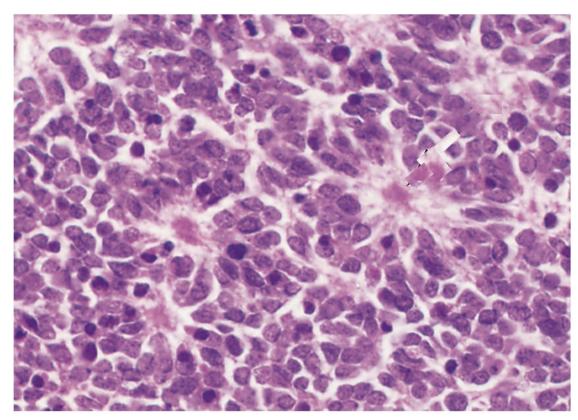


Another classic Rb finding (albeit neither as infamous nor as easy to spot as F-W rosettes) is depicted here—can you spot it? **Here** it is.

What's the finding?

Homer Wright rosettes in retinoblastoma (Rb). Not pathognomonic for Rb; they are found in other neuroblastic tumors.



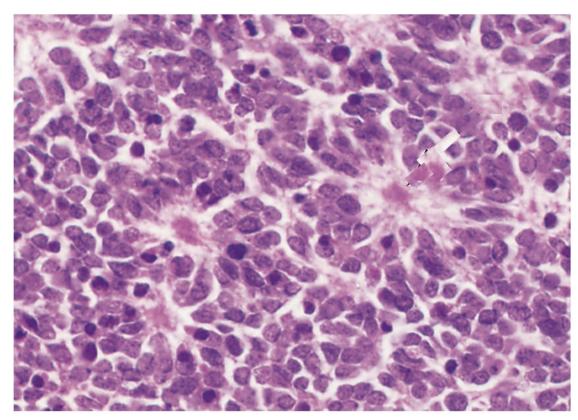


Another classic Rb finding (albeit neither as infamous nor as easy to spot as F-W rosettes) is depicted here—can you spot it? **Here** it is.

What's the finding?

Homer Wright rosettes in retinoblastoma (Rb). Not pathognomonic for Rb; they are found in other neuroblastic tumors. Also represents tumor process.



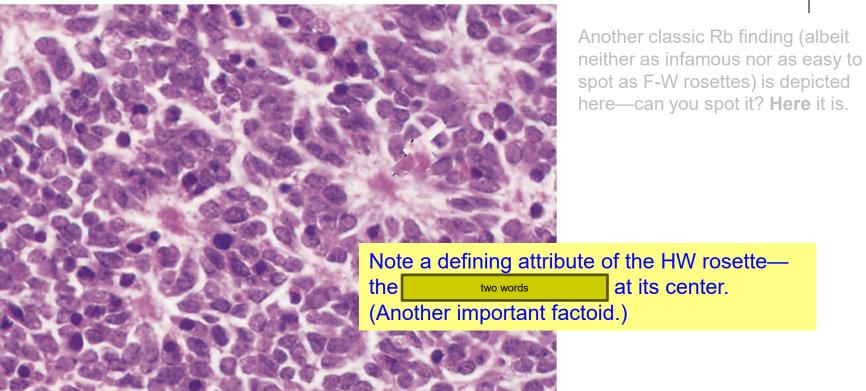


Another classic Rb finding (albeit neither as infamous nor as easy to spot as F-W rosettes) is depicted here—can you spot it? **Here** it is.

What's the finding?

Homer Wright rosettes in **retinoblastoma** (Rb). Not pathognomonic for Rb; they are found in other neuroblastic tumors. Also represents tumor differentiation.

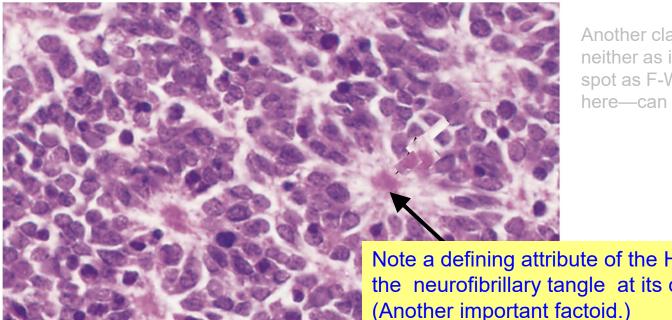




What's the finding?

Homer Wright rosettes in **retinoblastoma** (Rb). Not pathognomonic for Rb; they are found in other neuroblastic tumors. Also represents tumor differentiation.





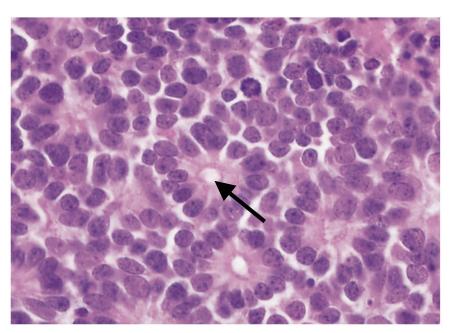
Another classic Rb finding (albeit neither as infamous nor as easy to spot as F-W rosettes) is depicted here—can you spot it? Here it is.

Note a defining attribute of the HW rosette the neurofibrillary tangle at its center. (Another important factoid.)

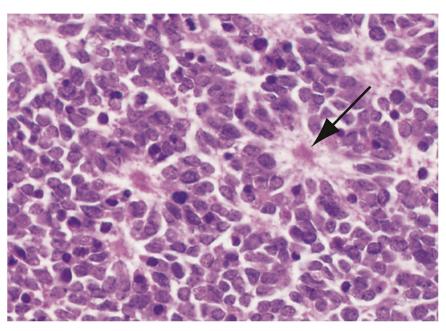
What's the finding?

Homer Wright rosettes in retinoblastoma (Rb). Not pathognomonic for Rb; they are found in other neuroblastic tumors. Also represents tumor differentiation.





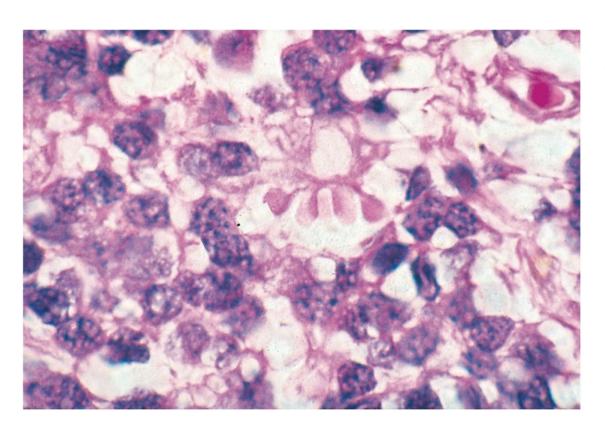
F-W rosette: Empty central lumen



HW rosette: Neurofibrillary tangle centrally

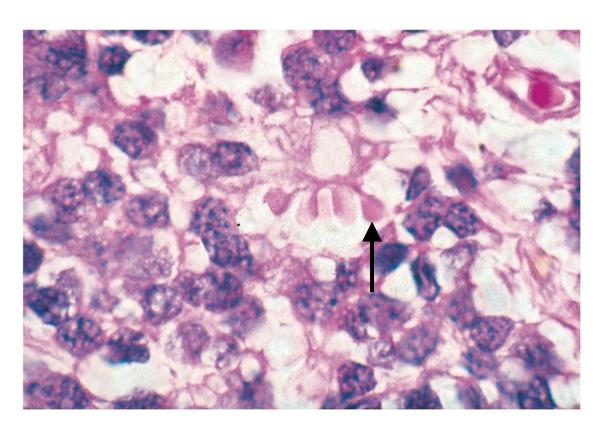
Flexner-Wintersteiner vs Homer Wright rosettes





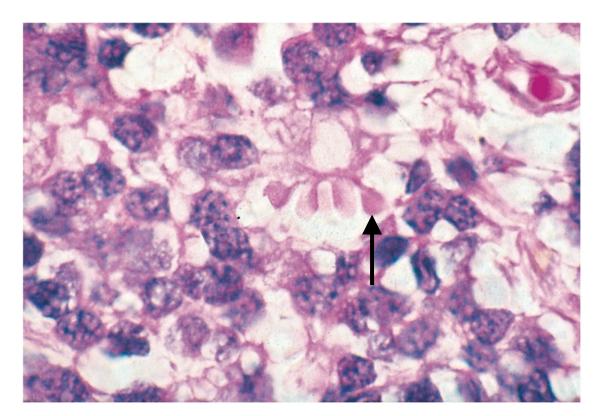
A third classic finding Rb, less infamous and spot-able still—see it?





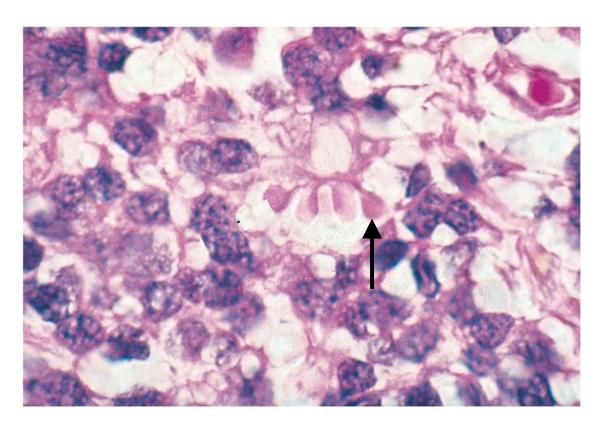
A third classic finding Rb, less infamous and spot-able still—see it? **Here** it is.





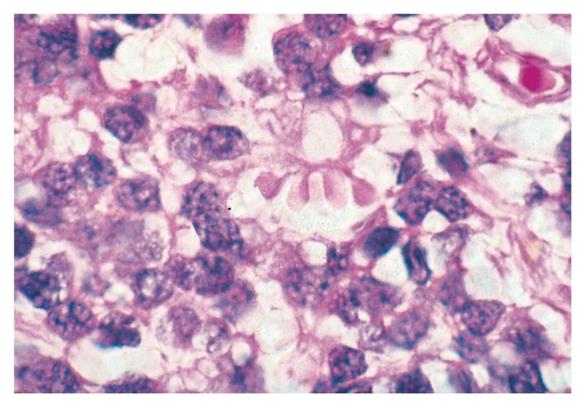
A third classic finding Rb, less infamous and spot-able still—see it? **Here** it is. The classic descriptor of this finding's shape is 'buzzterm'.'





A third classic finding Rb, less infamous and spot-able still—see it? **Here** it is. The classic descriptor of this finding's shape is 'bulbous.'

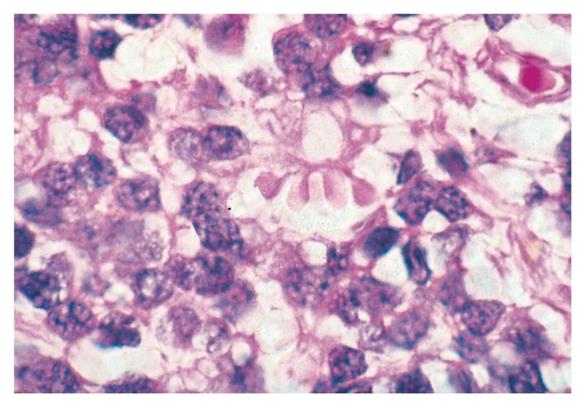




What's the finding?

A third classic finding Rb, less infamous and spot-able still—see it? **Here** it is. The classic descriptor of this finding's shape is 'bulbous.'



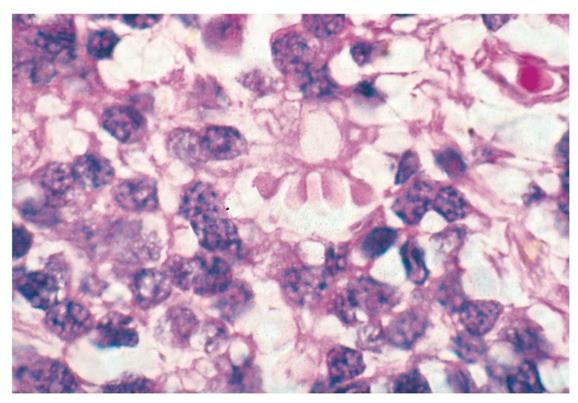


A third classic finding Rb, less infamous and spot-able still—see it? **Here** it is. The classic descriptor of this finding's shape is 'bulbous.'

What's the finding?

in **retinoblastoma** (Rb) .



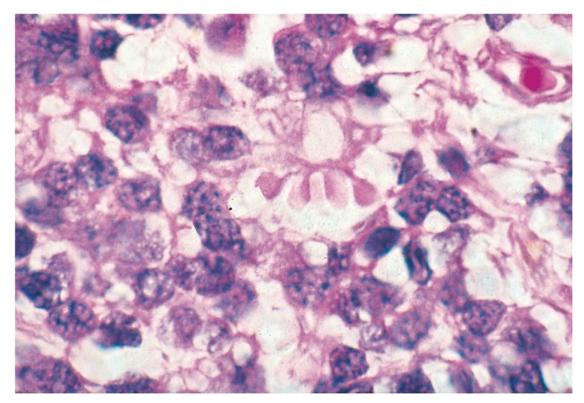


A third classic finding Rb, less infamous and spot-able still—see it? **Here** it is. The classic descriptor of this finding's shape is 'bulbous.'

What's the finding?

Fleurettes in retinoblastoma (Rb).



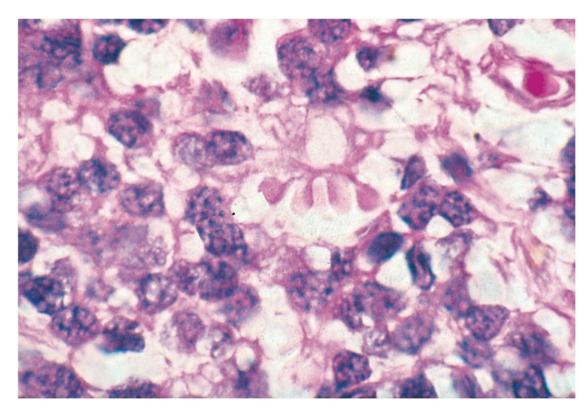


A third classic finding Rb, less infamous and spot-able still—see it? **Here** it is. The classic descriptor of this finding's shape is 'bulbous.'

What's the finding?

Fleurettes in retinoblastoma (Rb). Also represents tumor differentiation, specifically differentiation.



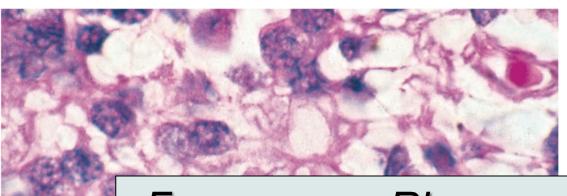


A third classic finding Rb, less infamous and spot-able still—see it? **Here** it is. The classic descriptor of this finding's shape is 'bulbous .'

What's the finding?

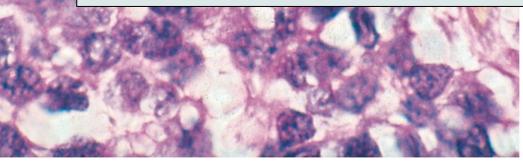
Fleurettes in **retinoblastoma** (Rb) . Also represents tumor differentiation, specifically photoreceptor differentiation. .





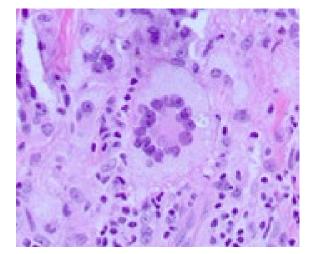
A third classic finding Rb, less infamous and spot-able still—see it? **Here** it is. The classic descriptor of this finding's shape is 'bulbous.'

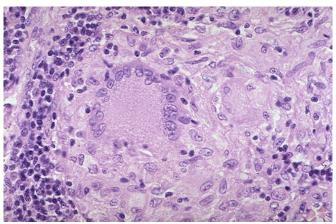
For more on Rb, see slide-set R2

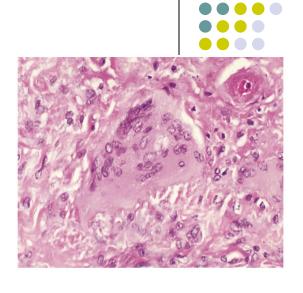


What's the finding?

Fleurettes in **retinoblastoma** (Rb) . Also represents tumor differentiation, specifically photoreceptor differentiation. .

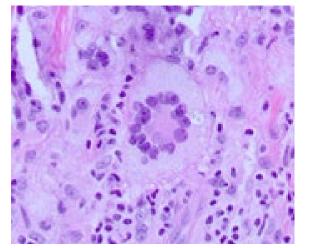


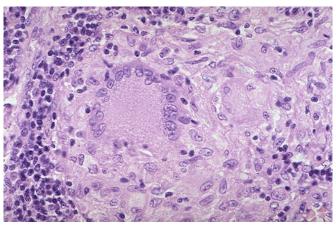


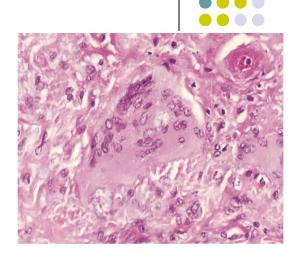


883

Another set of classic findings that must become readily recognizable.

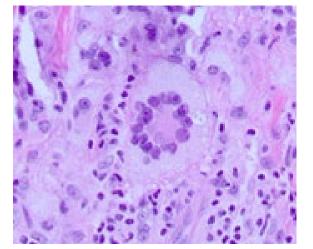


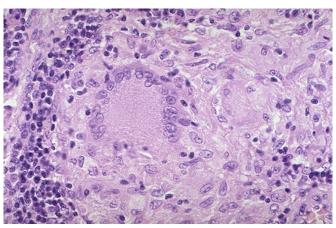


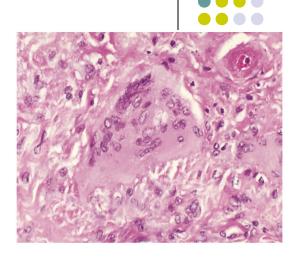


884

Another set of classic findings that must become readily recognizable. First things first—in general terms, what does each image depict? A two words

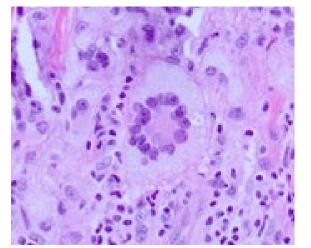


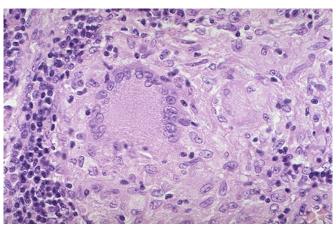


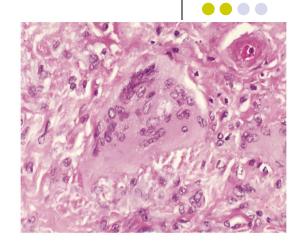


885

Another set of classic findings that must become readily recognizable. First things first—in general terms, what does each image depict? A *giant cell*

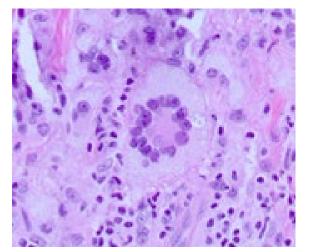


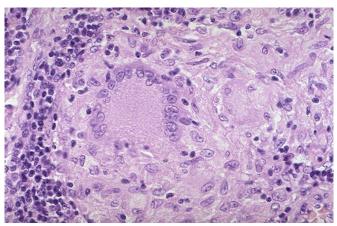


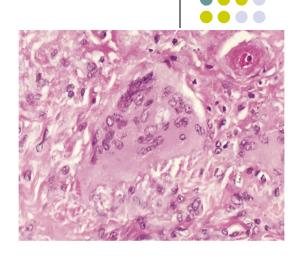


886

Another set of classic findings that must become readily recognizable. First things first—in general terms, what does each image depict? A *giant cell*, which is just a fancy word of two names for one cell type.

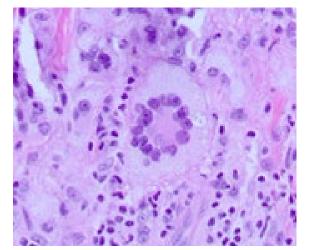


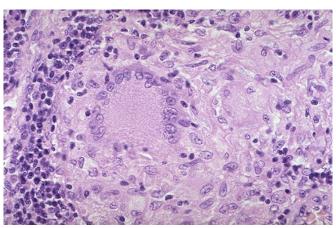


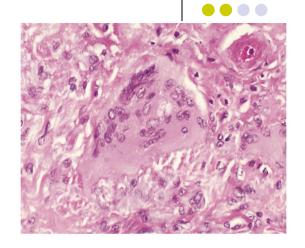


887

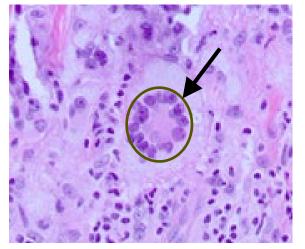
Another set of classic findings that must become readily recognizable. First things first—in general terms, what does each image depict? A *giant cell*, which is just a syncytium of histiocytes/macrophages.

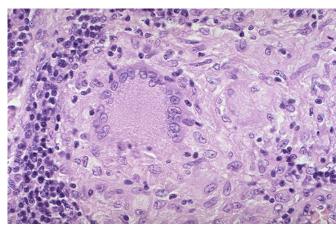


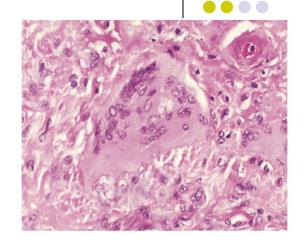




888





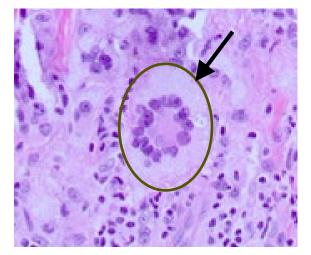


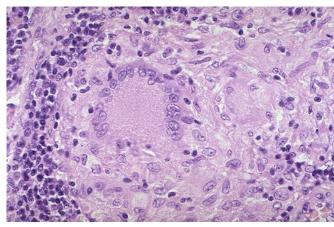
889

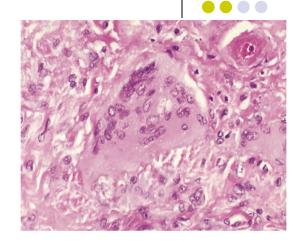
Note:

--Crucially: *This* is **not** the giant

cell...





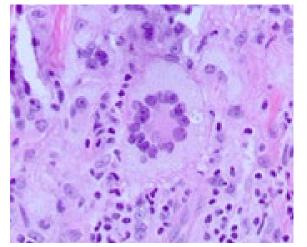


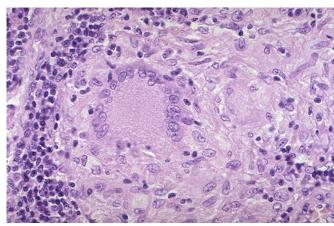
890

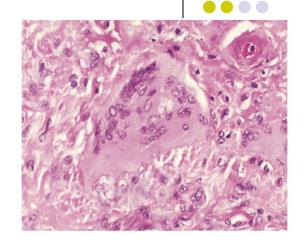
Note:

--Crucially: *This* is **not** the giant

cell...*This* is







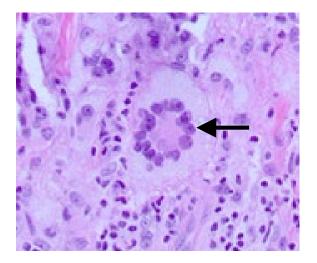
891

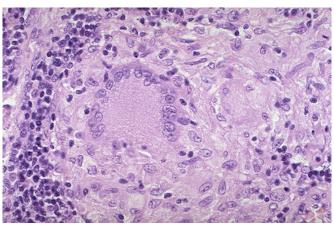
Note:

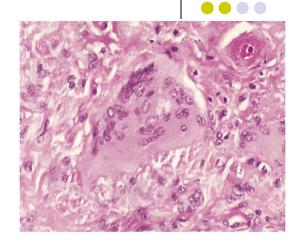
--Crucially: *This* is **not** the giant

cell...This is

-- location ring of nuclei







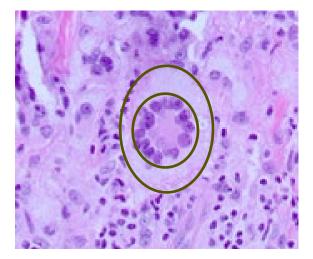
892

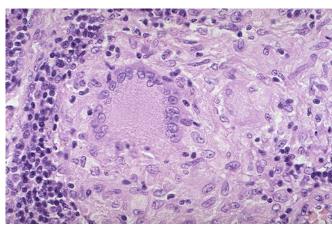
Note:

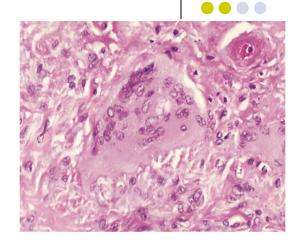
--Crucially: *This* is **not** the giant

cell...*This* is

--Central ring of nuclei







893

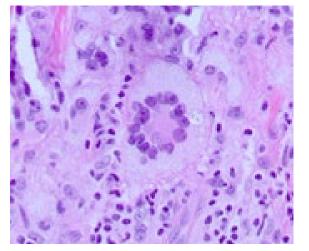
Note:

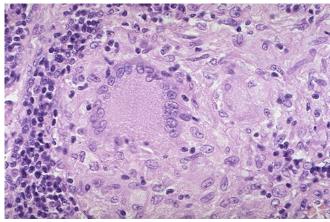
--Crucially: *This* is **not** the giant

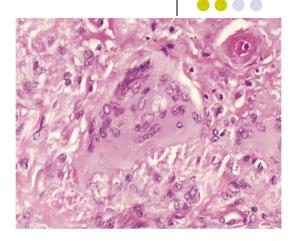
cell... This is

--Central ring of nuclei

--A surrounding donut of lipid







894

? giant cell

Note:

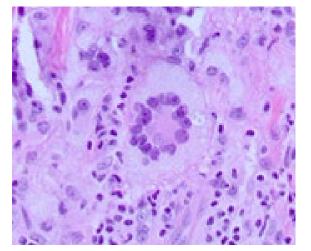
--Crucially: *This* is **not** the giant

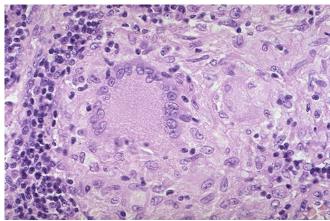
cell... This is

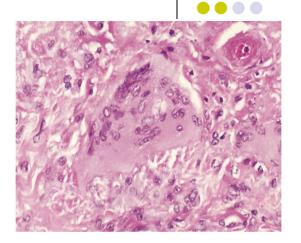
- --Central ring of nuclei
- --A surrounding donut of lipid

Another set of classic findings that must become readily recognizable. First things first—in general terms, what does each image depict? A *giant cell*, which is just a syncytium of histiocytes/macrophages. Now comes the challenging part—IDing each giant-cell type. Let's go through them.

What's the finding?







895

Touton giant cell

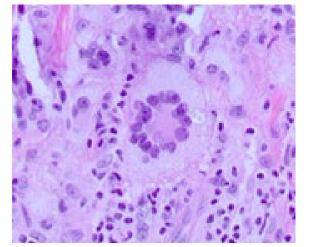
Note:

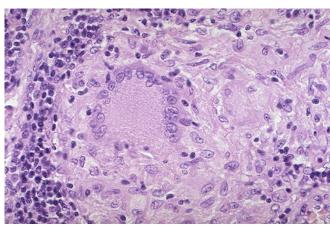
--Crucially: *This* is **not** the giant cell... *This* is

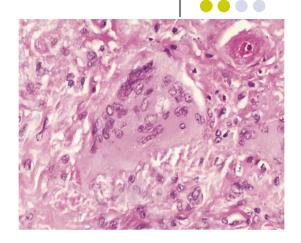
- --Central ring of nuclei
- --A surrounding donut of lipid

Another set of classic findings that must become readily recognizable. First things first—in general terms, what does each image depict? A *giant cell*, which is just a syncytium of histiocytes/macrophages. Now comes the challenging part—IDing each giant-cell type. Let's go through them.

What's the finding?







896

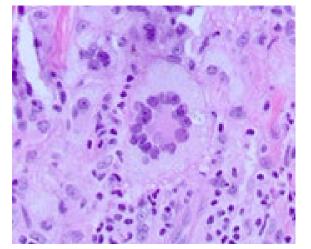
Touton giant cell

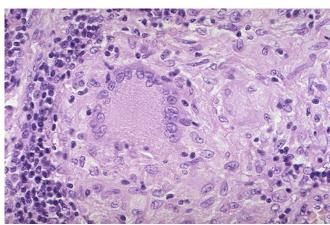
Touton giant cells are most closely associated with

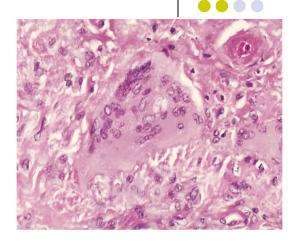
two words

-- Ochiral Ting of Huciel

--A surrounding donut of lipid





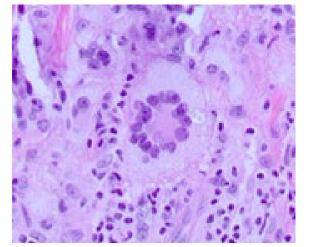


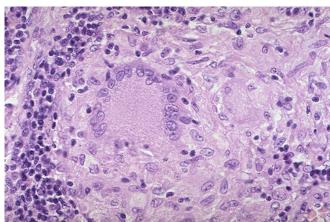
897

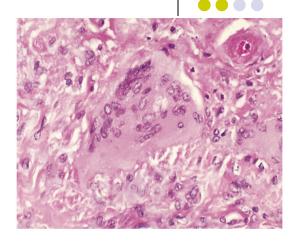
Touton giant cell

Touton giant cells are most closely associated with juvenile xanthogranuloma (JXG).

- -- Ochtrar Ting of Hucier
- --A surrounding donut of lipid





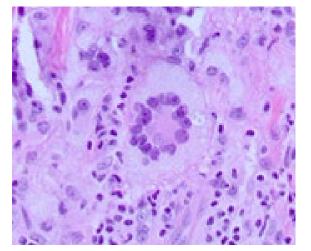


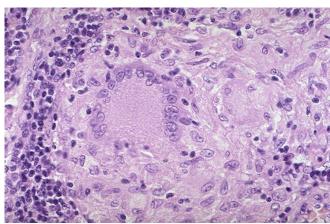
898

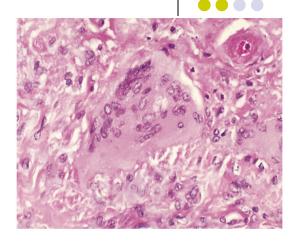
Touton giant cell

Touton giant cells are most closely associated with juvenile xanthogranuloma (JXG). However, they are also associated with two words xanthogranuloma dz.

--A surrounding donut of lipid







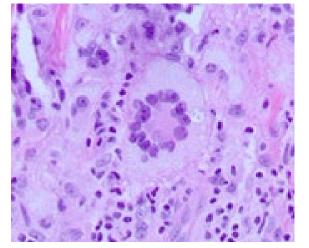
899

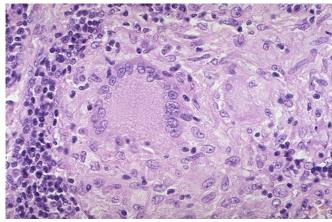
Touton giant cell

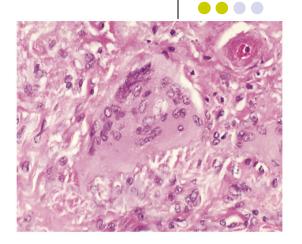
Touton giant cells are most closely associated with juvenile xanthogranuloma (JXG). However, they are also associated with adult-onset xanthogranuloma dz.

-- Ochilar hing of hacier

--A surrounding donut of lipid







900

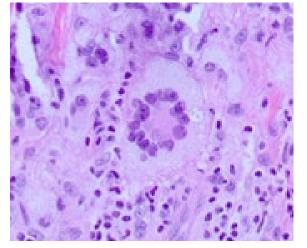
Touton giant cell

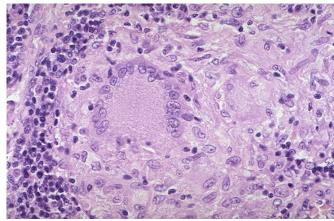
Note:

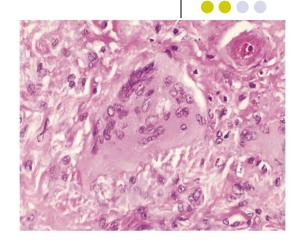
- --Crucially: *This* is **not** the giant cell... This is
- --Central ring of nuclei
- --A surrounding donut of lipid

Note: location

ring of nuclei







901

Touton giant cell

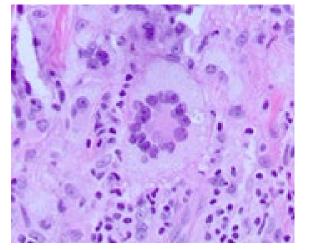
Note:

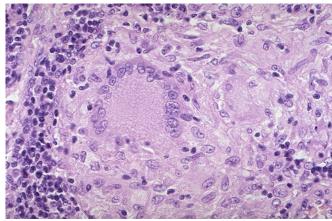
--Crucially: *This* is **not** the giant cell... *This* is

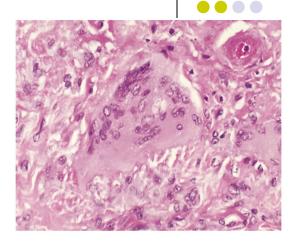
- --Central ring of nuclei
- --A surrounding donut of lipid

Note:

--Peripheral ring of nuclei







902

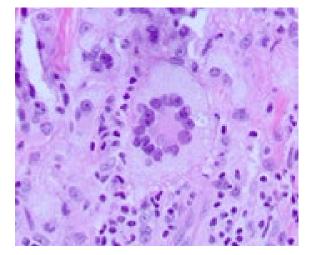
Touton giant cell

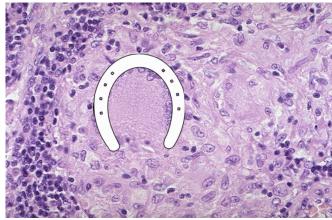
Note:

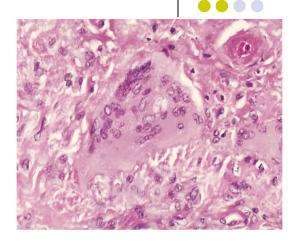
- --Crucially: *This* is **not** the giant cell... *This* is
- --Central ring of nuclei
- --A surrounding donut of lipid

Note:

--Peripheral ring of nuclei (often in a shape)







903

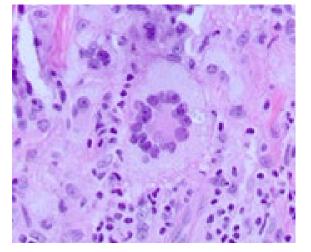
Touton giant cell

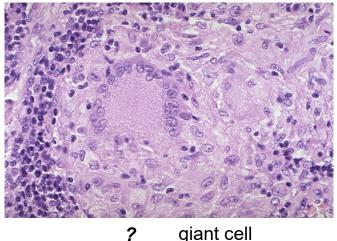
Note:

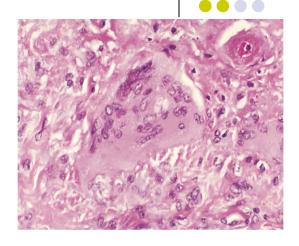
- --Crucially: *This* is **not** the giant cell... *This* is
- --Central ring of nuclei
- --A surrounding donut of lipid

Note:

--Peripheral ring of nuclei (often in a horseshoe)







904

Touton giant cell

Note:

--Crucially: *This* is **not** the giant cell... *This* is

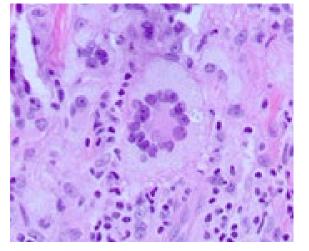
- --Central ring of nuclei
- --A surrounding donut of lipid

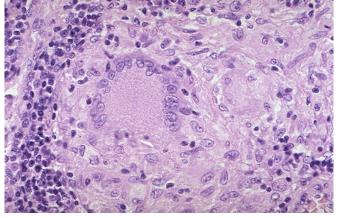
Note:

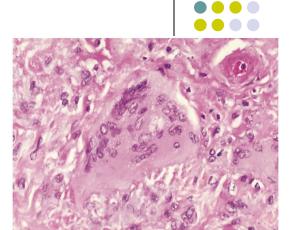
--Peripheral ring of nuclei (often in a horseshoe)

Another set of classic findings that must become readily recognizable. First things first—in general terms, what does each image depict? A *giant cell*, which is just a syncytium of histiocytes/macrophages. Now comes the challenging part—IDing each giant-cell type. Let's go through them.

What's the finding?







905

Touton giant cell

Langhans giant cell

Note:

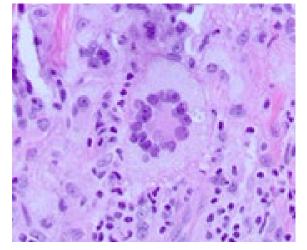
- --Crucially: *This* is **not** the giant cell... *This* is
- --Central ring of nuclei
- --A surrounding donut of lipid

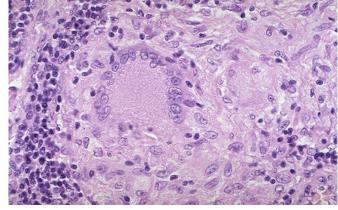
Note:

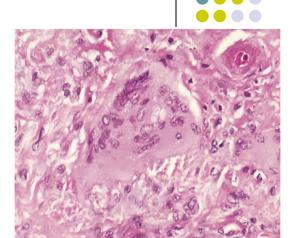
--Peripheral ring of nuclei (often in a horseshoe)

Another set of classic findings that must become readily recognizable. First things first—in general terms, what does each image depict? A *giant cell*, which is just a syncytium of histiocytes/macrophages. Now comes the challenging part—IDing each giant-cell type. Let's go through them.

What's the finding?







906

Touton giant cell

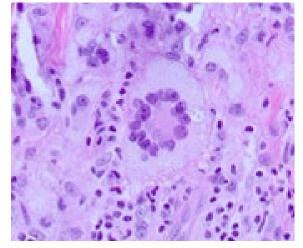
Note:

--Crucially: *This* is **not** the giant cell... *This* is

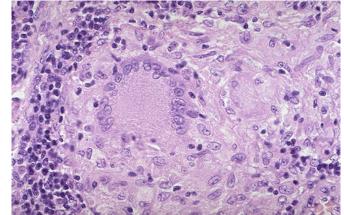
- --Central ring of nuclei
- --A surrounding donut of lipid

Langhans giant cell

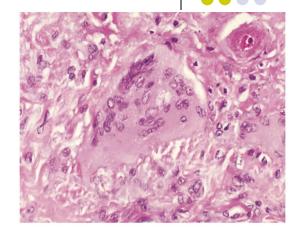
Langhans giant cells are associated with infectious etiologies, one of particular note being abb.







Langhans giant cell

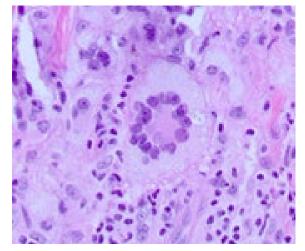


907

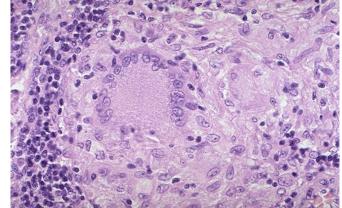
Note:

- --Crucially: *This* is **not** the giant cell... *This* is
- --Central ring of nuclei
- --A surrounding donut of lipid

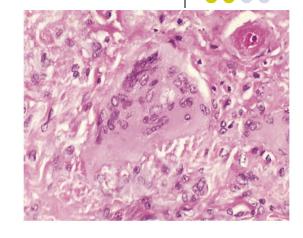
Langhans giant cells are associated with infectious etiologies, one of particular note being TB







Langhans giant cell



Note:

- --Crucially: *This* is **not** the giant cell... This is
- -- Central ring of nuclei
- --A surrounding donut of lipid

Note:

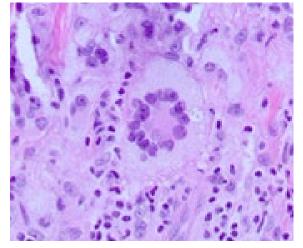
--Peripheral ring of nuclei (often in a horseshoe)

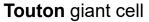
Note:

adjective nuclei

arrangement of

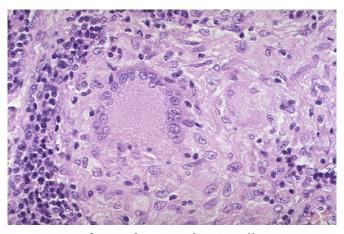
908







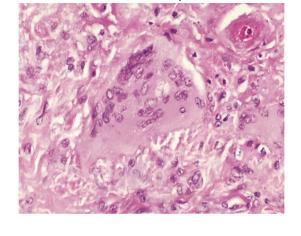
- --Crucially: *This* is **not** the giant cell... *This* is
- --Central ring of nuclei
- --A surrounding donut of lipid



Langhans giant cell

Note:

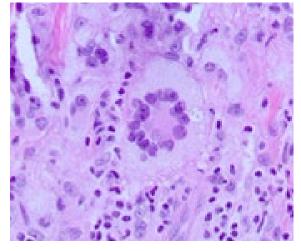
--Peripheral ring of nuclei (often in a horseshoe)



909

Note:

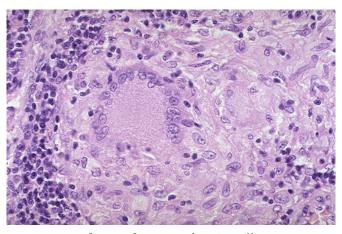
--Haphazard arrangement of nuclei



Touton giant cell



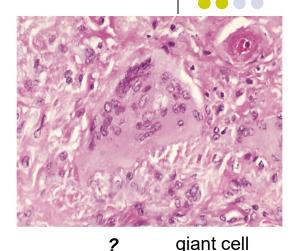
- --Crucially: *This* is **not** the giant cell... *This* is
- --Central ring of nuclei
- --A surrounding donut of lipid



Langhans giant cell

Note:

--Peripheral ring of nuclei (often in a horseshoe)



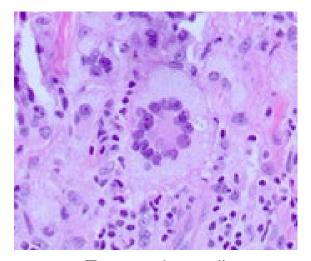
910

Note:

--Haphazard arrangement of nuclei

Another set of classic findings that must become readily recognizable. First things first—in general terms, what does each image depict? A *giant cell*, which is just a syncytium of histiocytes/macrophages. Now comes the challenging part—IDing each giant-cell type. Let's go through them.

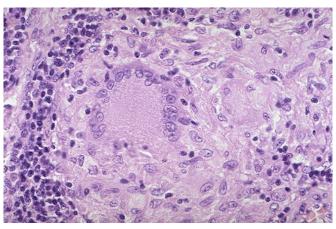
What's the finding?



Touton giant cell



- --Crucially: *This* is **not** the giant cell... *This* is
- --Central ring of nuclei
- --A surrounding donut of lipid



Langhans giant cell

Note:

--Peripheral ring of nuclei (often in a horseshoe)



911

Foreign-body giant cell

Note:

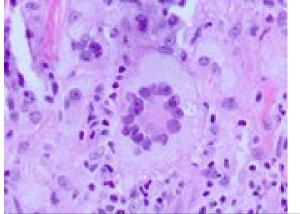
--Haphazard arrangement of nuclei

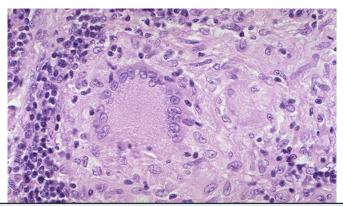
Another set of classic findings that must become readily recognizable. First things first—in general terms, what does each image depict? A *giant cell*, which is just a syncytium of histiocytes/macrophages. Now comes the challenging part—IDing each giant-cell type. Let's go through them.

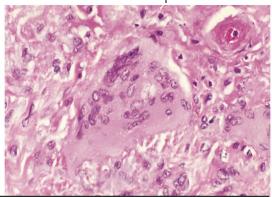
What's the finding?











For more on giant cells, see slide-set K20

Note:

- -- Crucially: *This* is **not** the giant cell... This is
- -- Central ring of nuclei
- -- A surrounding donut of lipid

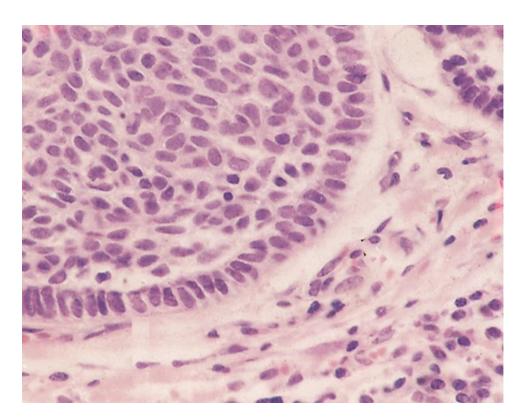
Note:

--Peripheral ring of nuclei (often in a horseshoe)

Note:

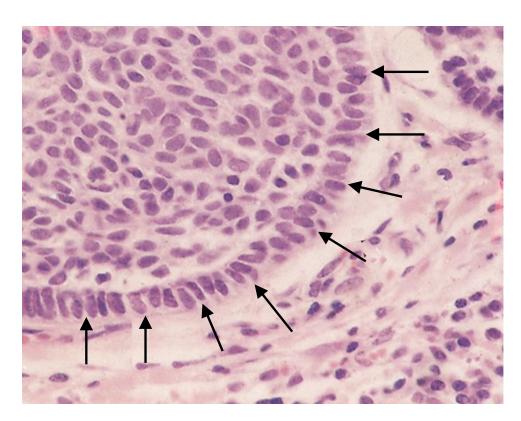
-- Haphazard arrangement of nuclei





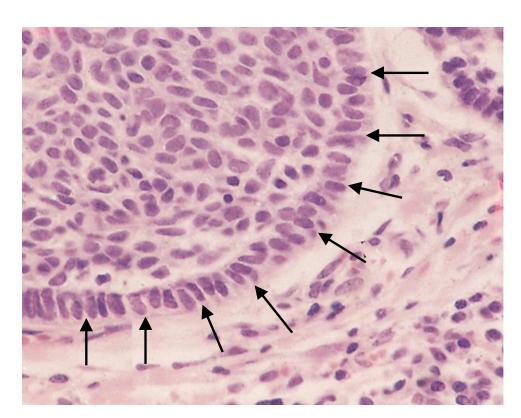
Another classic Path finding—what is it?





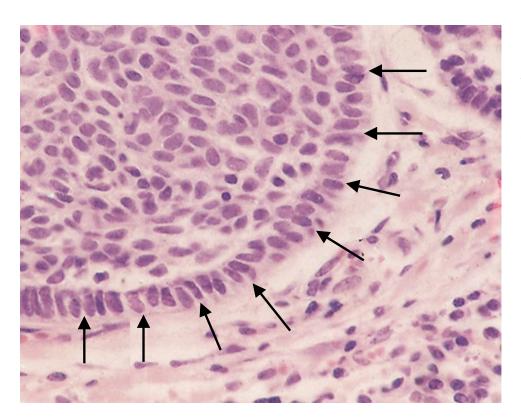
Another classic Path finding—what is it? It's **this** 'picket fence' of cells (*arrows*) on the border of a large nest of similar-appearing cells.





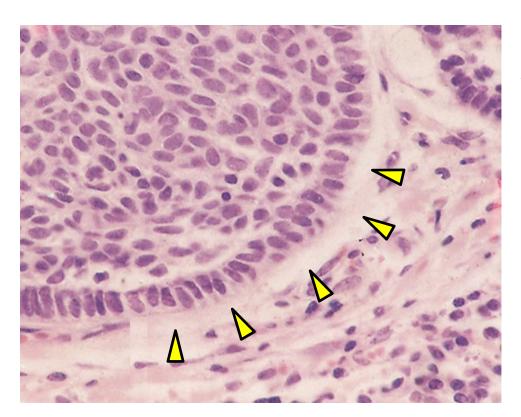
Another classic Path finding—what is it? It's **this** 'picket fence' of cells (*arrows*) on the border of a large nest of similar-appearing cells. In Path-speak the cells are said to be





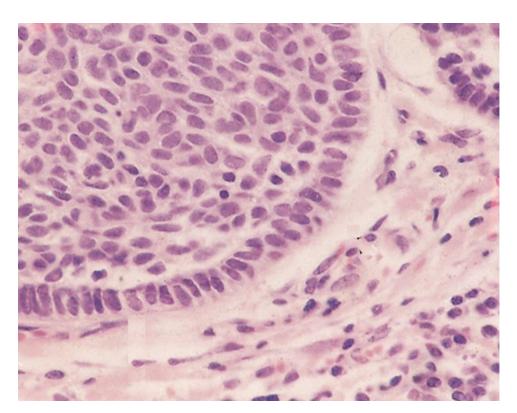
Another classic Path finding—what is it? It's **this** 'picket fence' of cells (*arrows*) on the border of a large nest of similar-appearing cells. In Path-speak the cells are said to be 'palisading'.





Another classic Path finding—what is it? It's **this** 'picket fence' of cells (*arrows*) on the border of a large nest of similar-appearing cells. In Path-speak the cells are said to be 'palisading'. Note also the clear space (*arrowheads*) between the palisading cells and the surrounding tissue.



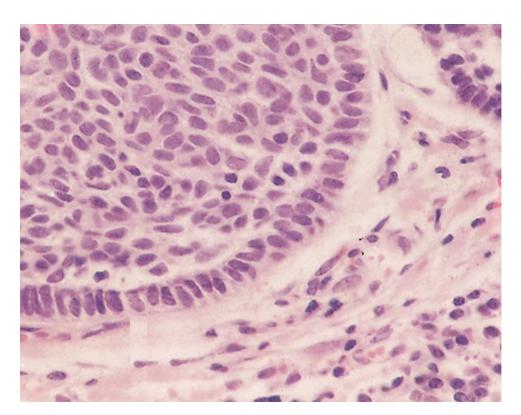


What's the diagnosis?

Another classic Path finding—what is it? It's **this** 'picket fence' of cells (*arrows*) on the border of a large nest of similar-appearing cells. In Path-speak the cells are said to be 'palisading'. Note also the clear space (*arrowheads*) between the palisading cells and the surrounding tissue.

When you see palisading cells with a surrounding empty cleft like this, one diagnosis should come to mind:





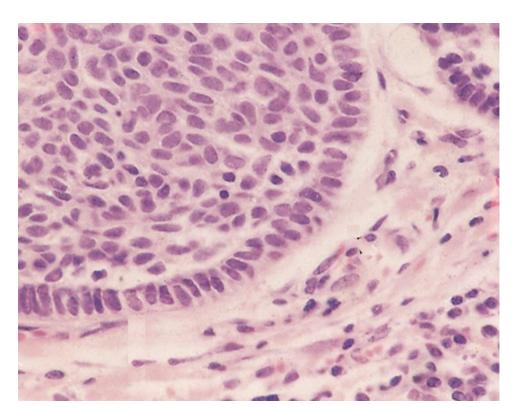
Another classic Path finding—what is it? It's this 'picket fence' of cells (arrows) on the border of a large nest of similar-appearing cells. In Path-speak the cells are said to be 'palisading'. Note also the clear space (arrowheads) between the palisading cells and the surrounding tissue.

When you see palisading cells with a surrounding empty cleft like this, one diagnosis should come to mind:

What's the diagnosis?

Basal cell carcinoma (BCC) is the most common malignancy of the structure





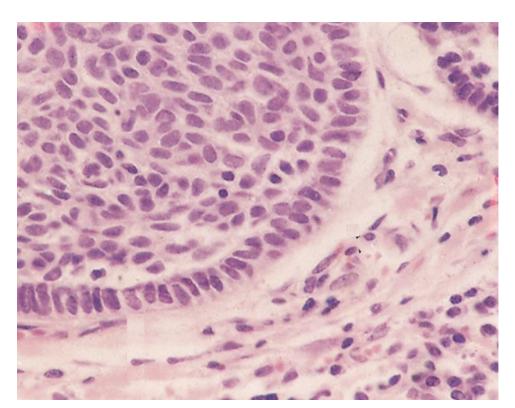
Another classic Path finding—what is it? It's **this** 'picket fence' of cells (*arrows*) on the border of a large nest of similar-appearing cells. In Path-speak the cells are said to be 'palisading'. Note also the clear space (*arrowheads*) between the palisading cells and the surrounding tissue.

When you see palisading cells with a surrounding empty cleft like this, one diagnosis should come to mind:

What's the diagnosis?

Basal cell carcinoma (BCC) is the most common malignancy of the eyelids





Another classic Path finding—what is it? It's **this** 'picket fence' of cells (*arrows*) on the border of a large nest of similar-appearing cells. In Path-speak the cells are said to be 'palisading'. Note also the clear space (*arrowheads*) between the palisading cells and the surrounding tissue.

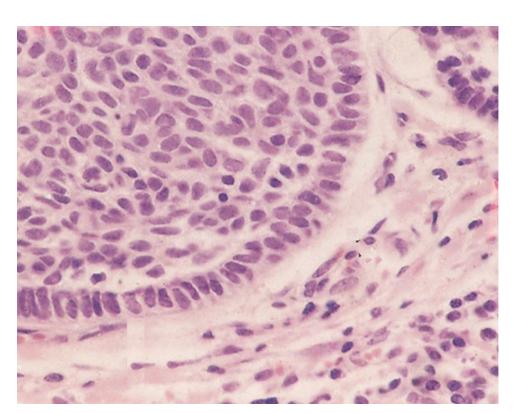
When you see palisading cells with a surrounding empty cleft like this, one diagnosis should come to mind:

What's the diagnosis?

Basal cell carcinoma (BCC) is the most common malignancy of the eyelids . is a strong risk factor

two words





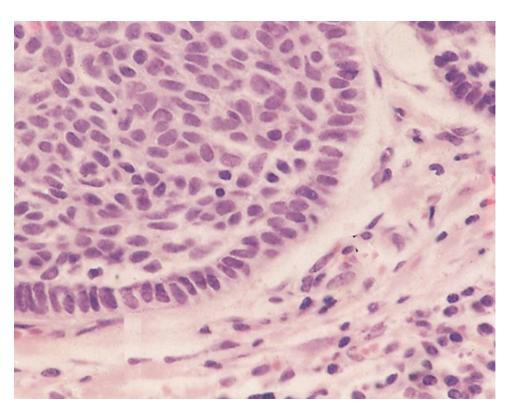
Another classic Path finding—what is it? It's **this** 'picket fence' of cells (*arrows*) on the border of a large nest of similar-appearing cells. In Path-speak the cells are said to be 'palisading'. Note also the clear space (*arrowheads*) between the palisading cells and the surrounding tissue.

When you see palisading cells with a surrounding empty cleft like this, one diagnosis should come to mind:

What's the diagnosis?

Basal cell carcinoma (BCC) is the most common malignancy of the eyelids . Sun exposure is a strong risk factor





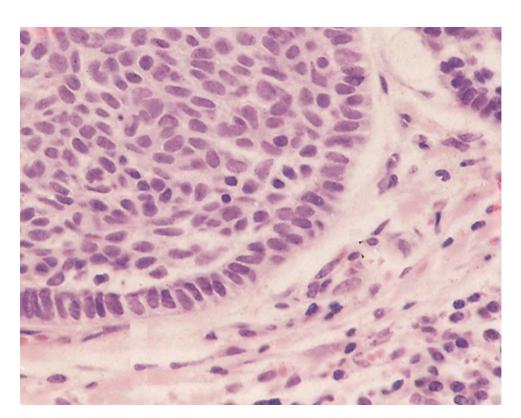
Another classic Path finding—what is it? It's **this** 'picket fence' of cells (*arrows*) on the border of a large nest of similar-appearing cells. In Path-speak the cells are said to be 'palisading'. Note also the clear space (*arrowheads*) between the palisading cells and the surrounding tissue.

When you see palisading cells with a surrounding empty cleft like this, one diagnosis should come to mind:

What's the diagnosis?

Basal cell carcinoma (BCC) is the most common malignancy of the eyelids. Sun exposure is a strong risk factor, and explains why the lower vs lid is more commonly affected.





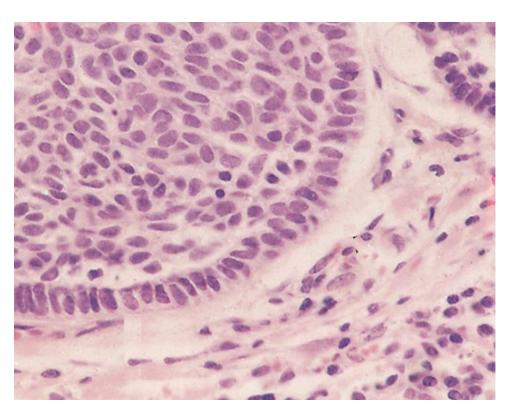
Another classic Path finding—what is it? It's **this** 'picket fence' of cells (*arrows*) on the border of a large nest of similar-appearing cells. In Path-speak the cells are said to be 'palisading'. Note also the clear space (*arrowheads*) between the palisading cells and the surrounding tissue.

When you see palisading cells with a surrounding empty cleft like this, one diagnosis should come to mind:

What's the diagnosis?

Basal cell carcinoma (BCC) is the most common malignancy of the eyelids. Sun exposure is a strong risk factor, and explains why the lower lid is more commonly affected.





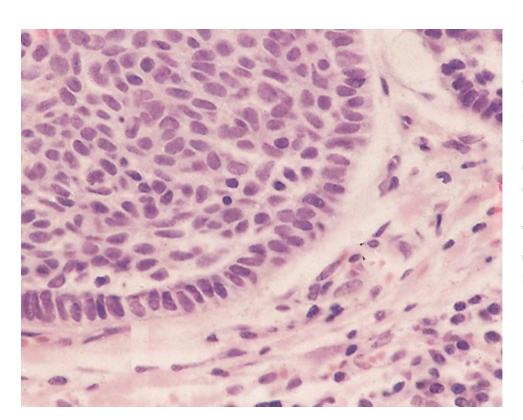
Another classic Path finding—what is it? It's **this** 'picket fence' of cells (*arrows*) on the border of a large nest of similar-appearing cells. In Path-speak the cells are said to be 'palisading'. Note also the clear space (*arrowheads*) between the palisading cells and the surrounding tissue.

When you see palisading cells with a surrounding empty cleft like this, one diagnosis should come to mind:

What's the diagnosis?

Basal cell carcinoma (BCC) is the most common malignancy of the eyelids . Sun exposure is a strong risk factor, and explains why the lower lid is more commonly affected. The clear spaces surrounding the tumor-cell islands are





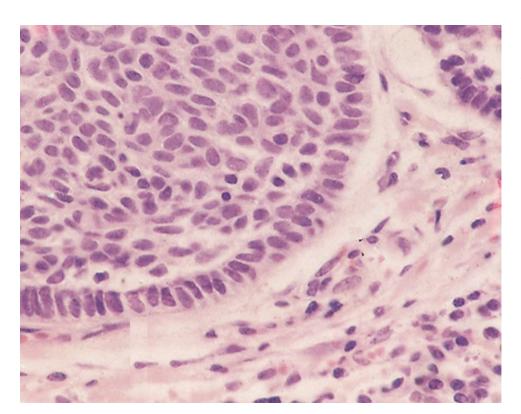
Another classic Path finding—what is it? It's **this** 'picket fence' of cells (*arrows*) on the border of a large nest of similar-appearing cells. In Path-speak the cells are said to be 'palisading'. Note also the clear space (*arrowheads*) between the palisading cells and the surrounding tissue.

When you see palisading cells with a surrounding empty cleft like this, one diagnosis should come to mind:

What's the diagnosis?

Basal cell carcinoma (BCC) is the most common malignancy of the eyelids. Sun exposure is a strong risk factor, and explains why the lower lid is more commonly affected. The clear spaces surrounding the tumor-cell islands are factitious





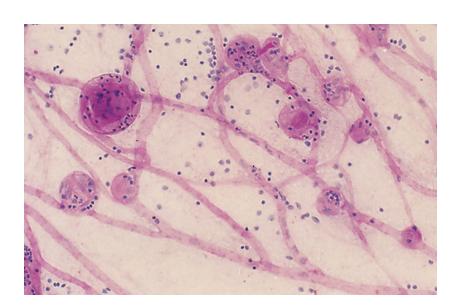
Another classic Path finding—what is it? It's **this** 'picket fence' of cells (*arrows*) on the border of a large nest of similar-appearing cells. In Path-speak the cells are said to be 'palisading'. Note also the clear space (*arrowheads*) between the palisading cells and the surrounding tissue.

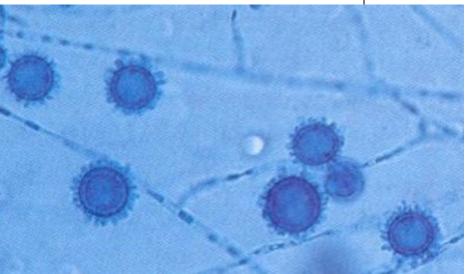
When you see palisading cells with a surrounding empty cleft like this, one diagnosis should come to mind:

What's the diagnosis?

Basal cell carcinoma (BCC) is the most common malignancy of the eyelids. Sun exposure is a strong risk factor, and explains why the lower lid is more commonly affected. The clear spaces surrounding the tumor-cell islands are factitious (they arise during tissue processing).

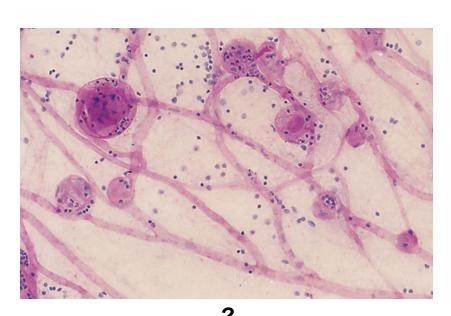


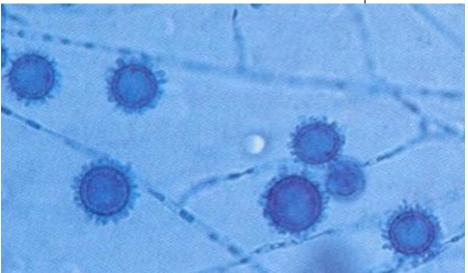




These pics could easily be confused for one another, but the path they depict couldn't be more different. What are they?



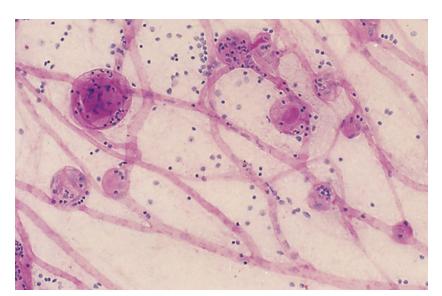


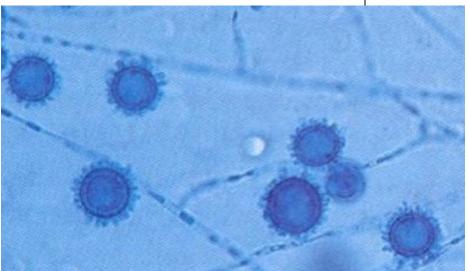


•

These pics could easily be confused for one another, but the path they depict couldn't be more different. What are they? The pic on the left is a special prep that allows one to see two words of the retinal vasculature.



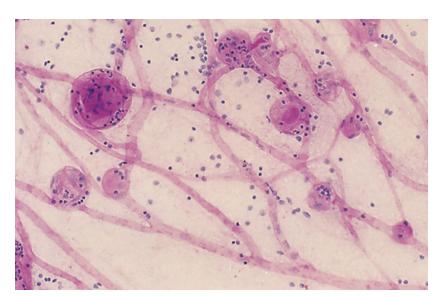


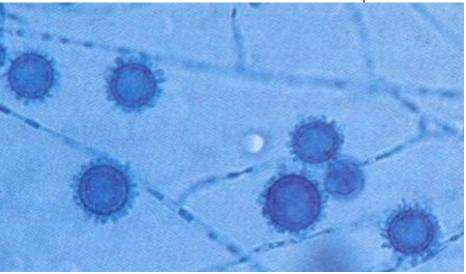


Diabetic microaneurysms (MA)

These pics could easily be confused for one another, but the path they depict couldn't be more different. What are they? The pic on the left is a special prep that allows one to see diabetic microaneurysms of the retinal vasculature.





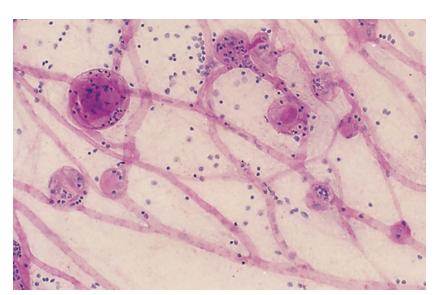


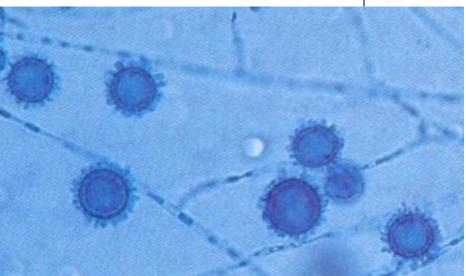
Diabetic microaneurysms (MA)

7

These pics could easily be confused for one another, but the path they depict couldn't be more different. What are they? The pic on the left is a special prep that allows one to see diabetic microaneurysms of the retinal vasculature. The pic on the right depicts...







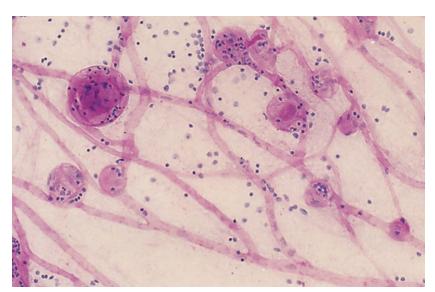
Diabetic microaneurysms (MA)

2

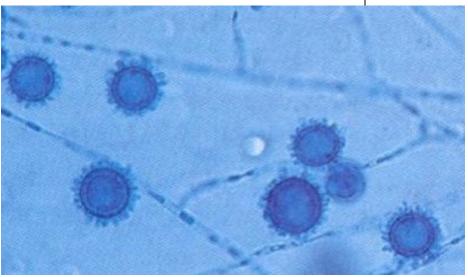
These pics could easily be confused for one another, but the path they depict couldn't be more different. What are they? The pic on the left is a special prep that allows one to see diabetic microaneurysms of the retinal vasculature. The pic on the right depicts...the filamentous/mold form of the dimorphic fungus

two words





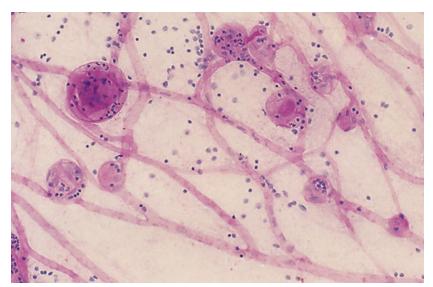




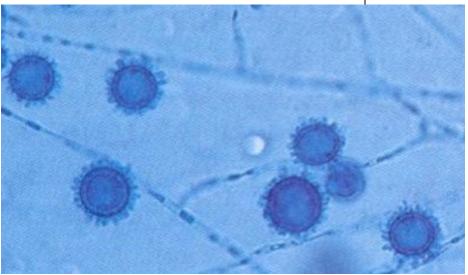
Histoplasma capsulatum

These pics could easily be confused for one another, but the path they depict couldn't be more different. What are they? The pic on the left is a special prep that allows one to see diabetic microaneurysms of the retinal vasculature. The pic on the right depicts...the filamentous/mold form of the dimorphic fungus *Histoplasma capsulatum*.







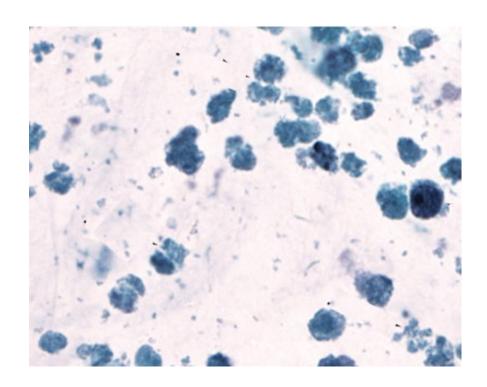


Histoplasma capsulatum

These pics could easily be confused for one another, but the path they depict couldn't be more different. What are they? The pic on the left is a special prep that allows one to see diabetic microaneurysms of the retinal vasculature. The pic on the right depicts...the filamentous/mold form of the dimorphic fungus *Histoplasma capsulatum*.

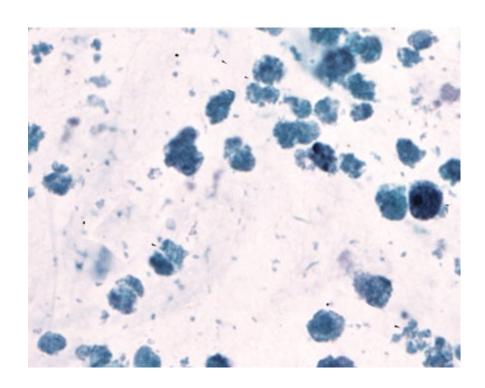
Take note of these strikingly similar images, lest you get fooled on an exam into thinking one is the other!





All you're told about this is it's a vitreous biopsy.

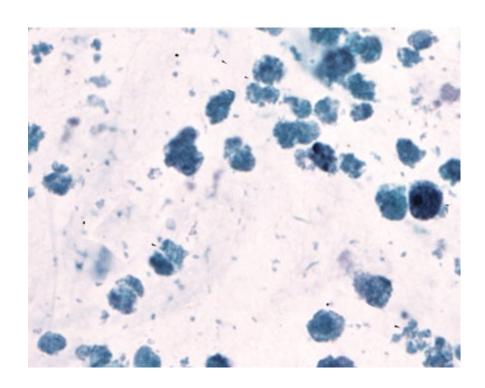




All you're told about this is it's a vitreous biopsy.

In three words, what do you see?

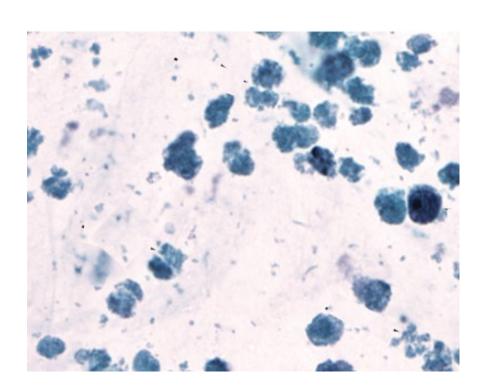




All you're told about this is it's a vitreous biopsy.

In three words, what do you see? Big blue cells

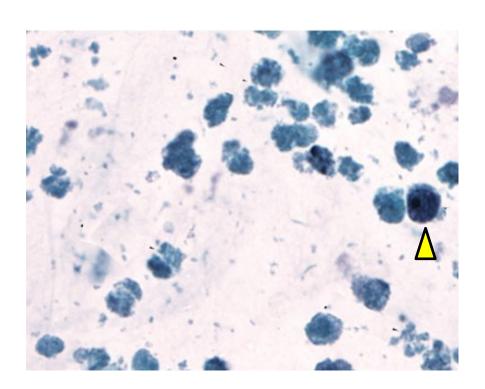




All you're told about this is it's a vitreous biopsy.

- --?
- --? --?





All you're told about this is it's a vitreous biopsy.

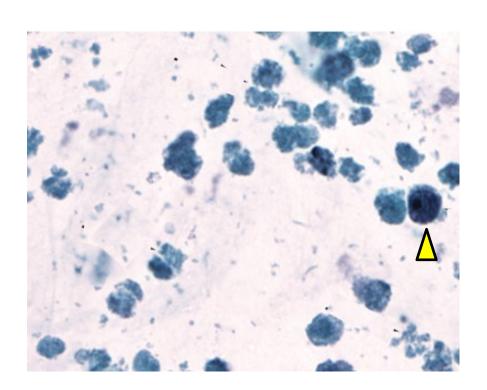
In three words, what do you see? Big blue cells
Note that these BBCs demonstrate several
worrisome characteristics:

--Scant (arrowhead)

--?

--?





All you're told about this is it's a vitreous biopsy.

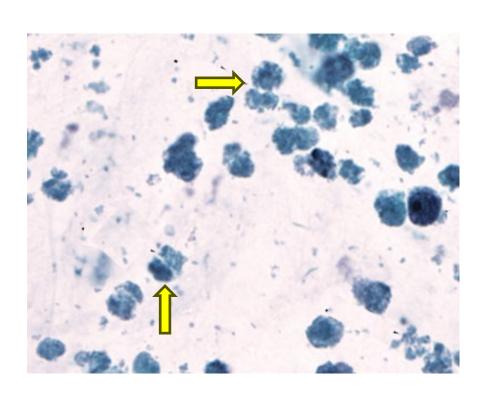
In three words, what do you see? Big blue cells
Note that these BBCs demonstrate several
worrisome characteristics:

--Scant cytoplasm (arrowhead)

--?

--?





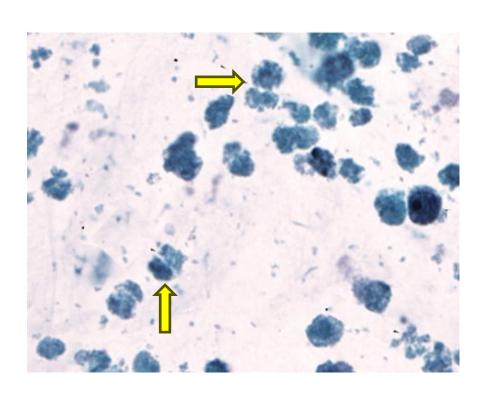
All you're told about this is it's a vitreous biopsy.

In three words, what do you see? Big blue cells
Note that these BBCs demonstrate several
worrisome characteristics:

--Scant cytoplasm (arrowhead)

-- cells (arrows)

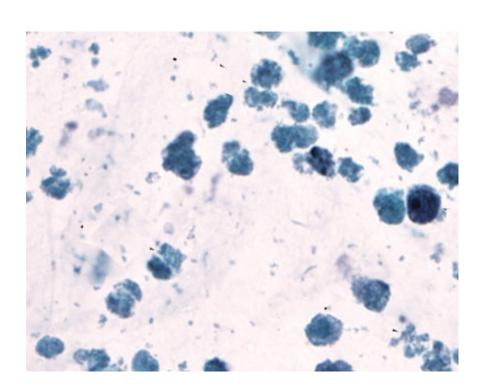




All you're told about this is it's a vitreous biopsy.

- --Scant cytoplasm (arrowhead)
- --Smudge cells (arrows)
- -- ?
- --?



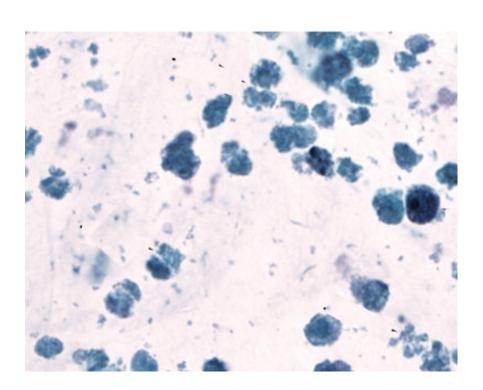


All you're told about this is it's a vitreous biopsy.

In three words, what do you see? Big blue cells
Note that these BBCs demonstrate several
worrisome characteristics:

- --Scant cytoplasm (arrowhead)
- --Smudge cells (arrows)
- --Hyperchromatic



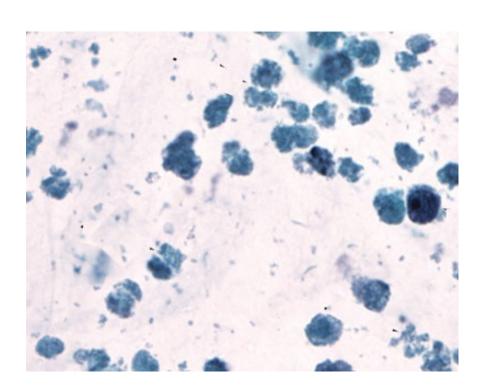


All you're told about this is it's a vitreous biopsy.

In three words, what do you see? Big blue cells
Note that these BBCs demonstrate several
worrisome characteristics:

- --Scant cytoplasm (arrowhead)
- --Smudge cells (arrows)
- --Hyperchromatic nuclei

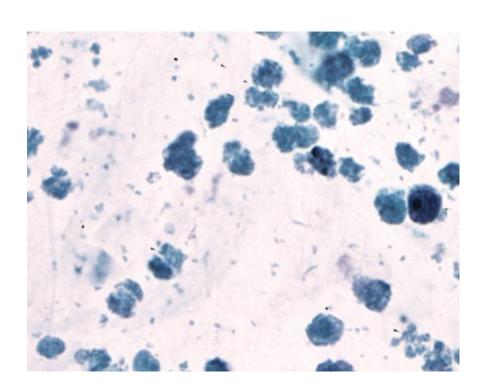




All you're told about this is it's a vitreous biopsy.

- --Scant cytoplasm (arrowhead)
- --Smudge cells (arrows)
- --Hyperchromatic nuclei
- --Prominent

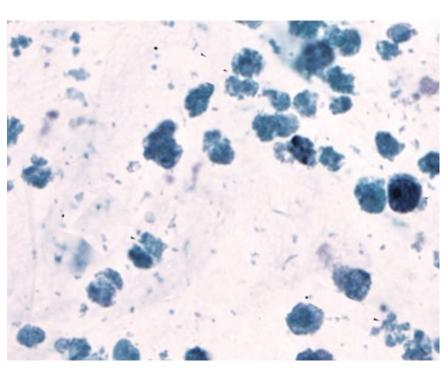




All you're told about this is it's a vitreous biopsy.

- --Scant cytoplasm (arrowhead)
- --Smudge cells (arrows)
- --Hyperchromatic nuclei
- --Prominent nucleoli





What's the diagnosis?

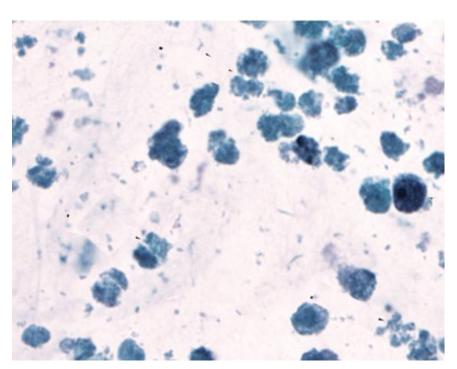
All you're told about this is it's a vitreous biopsy.

In three words, what do you see? Big blue cells
Note that these BBCs demonstrate several
worrisome characteristics:

- --Scant cytoplasm (arrowhead)
- --Smudge cells (arrows)
- --Hyperchromatic nuclei
- --Prominent nucleoli

When you see wonky BBCs on a vitreous biopsy, one dx should come to mind:





What's the diagnosis?

All you're told about this is it's a vitreous biopsy.

In three words, what do you see? Big blue cells Note that these BBCs demonstrate several worrisome characteristics:

- --Scant cytoplasm (arrowhead)
- --Smudge cells (arrows)
- --Hyperchromatic nuclei
- --Prominent nucleoli

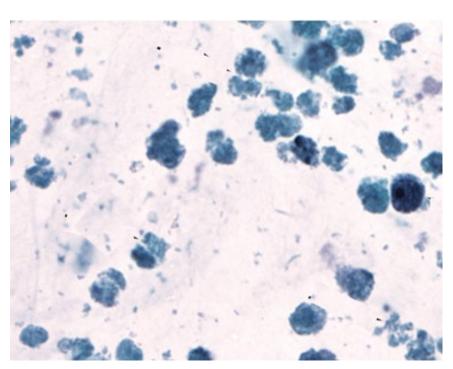
When you see wonky BBCs on a vitreous biopsy, one dx should come to mind:

Primary intraocular lymphoma (PIOL) is



malignancy





What's the diagnosis?

All you're told about this is it's a vitreous biopsy.

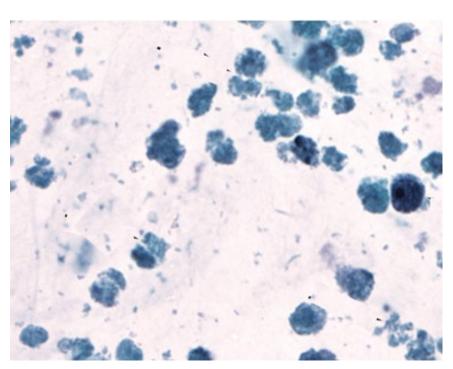
In three words, what do you see? Big blue cells Note that these BBCs demonstrate several worrisome characteristics:

- --Scant cytoplasm (arrowhead)
- --Smudge cells (arrows)
- --Hyperchromatic nuclei
- --Prominent nucleoli

When you see wonky BBCs on a vitreous biopsy, one dx should come to mind:

Primary intraocular lymphoma (PIOL) is an uncommon malignancy





What's the diagnosis?

All you're told about this is it's a vitreous biopsy.

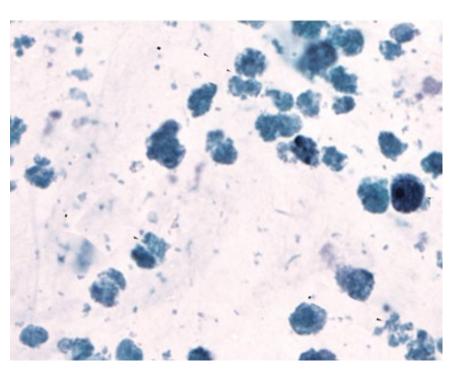
In three words, what do you see? Big blue cells
Note that these BBCs demonstrate several
worrisome characteristics:

- --Scant cytoplasm (arrowhead)
- --Smudge cells (arrows)
- --Hyperchromatic nuclei
- --Prominent nucleoli

When you see wonky BBCs on a vitreous biopsy, one dx should come to mind:

Primary intraocular lymphoma (PIOL) is an uncommon malignancy. The vast majority are in origin.





What's the diagnosis?

All you're told about this is it's a vitreous biopsy.

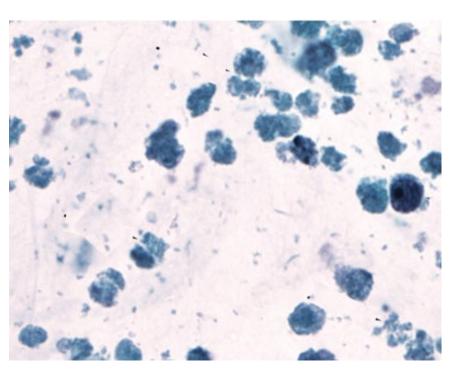
In three words, what do you see? Big blue cells
Note that these BBCs demonstrate several
worrisome characteristics:

- --Scant cytoplasm (arrowhead)
- --Smudge cells (arrows)
- --Hyperchromatic nuclei
- --Prominent nucleoli

When you see wonky BBCs on a vitreous biopsy, one dx should come to mind:

Primary intraocular lymphoma (PIOL) is an uncommon malignancy. The vast majority are B-cell in origin.





What's the diagnosis?

All you're told about this is it's a vitreous biopsy.

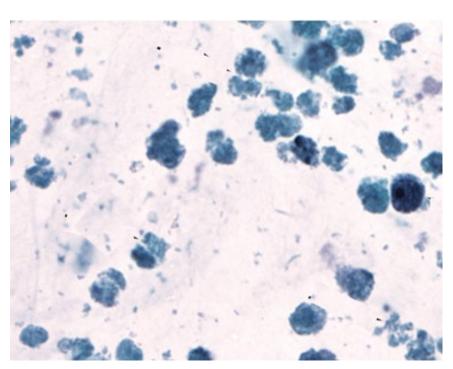
In three words, what do you see? Big blue cells Note that these BBCs demonstrate several worrisome characteristics:

- --Scant cytoplasm (arrowhead)
- --Smudge cells (arrows)
- --Hyperchromatic nuclei
- --Prominent nucleoli

When you see wonky BBCs on a vitreous biopsy, one dx should come to mind:

Primary intraocular lymphoma (PIOL) is an uncommon malignancy. The vast majority are B-cell in origin. At least half of pts demonstrate abb. involvement.





What's the diagnosis?

All you're told about this is it's a vitreous biopsy.

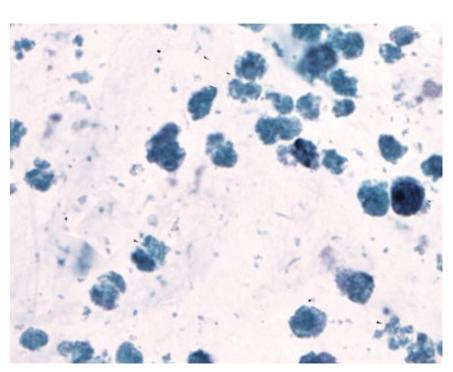
In three words, what do you see? Big blue cells
Note that these BBCs demonstrate several
worrisome characteristics:

- --Scant cytoplasm (arrowhead)
- --Smudge cells (arrows)
- --Hyperchromatic nuclei
- --Prominent nucleoli

When you see wonky BBCs on a vitreous biopsy, one dx should come to mind:

Primary intraocular lymphoma (PIOL) is an uncommon malignancy. The vast majority are B-cell in origin. At least half of pts demonstrate CNS involvement.





What's the diagnosis?

All you're told about this is it's a vitreous biopsy.

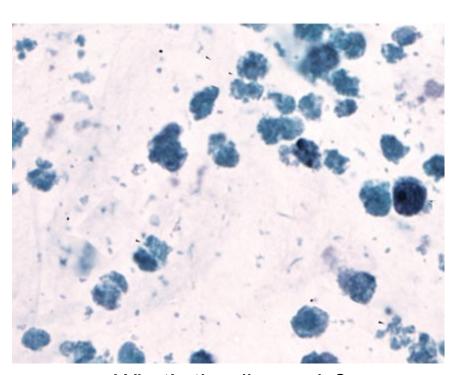
In three words, what do you see? Big blue cells
Note that these BBCs demonstrate several
worrisome characteristics:

- --Scant cytoplasm (arrowhead)
- --Smudge cells (arrows)
- --Hyperchromatic nuclei
- --Prominent nucleoli

When you see wonky BBCs on a vitreous biopsy, one dx should come to mind:

Primary intraocular lymphoma (PIOL) is an uncommon malignancy. The vast majority are B-cell in origin. At least half of pts demonstrate CNS involvement. The prognosis is





What's the diagnosis?

All you're told about this is it's a vitreous biopsy.

In three words, what do you see? Big blue cells Note that these BBCs demonstrate several worrisome characteristics:

- --Scant cytoplasm (arrowhead)
- --Smudge cells (arrows)
- --Hyperchromatic nuclei
- --Prominent nucleoli

When you see wonky BBCs on a vitreous biopsy, one dx should come to mind:

Primary intraocular lymphoma (PIOL) is an uncommon malignancy. The vast majority are B-cell in origin. At least half of pts demonstrate CNS involvement. The prognosis is poor.