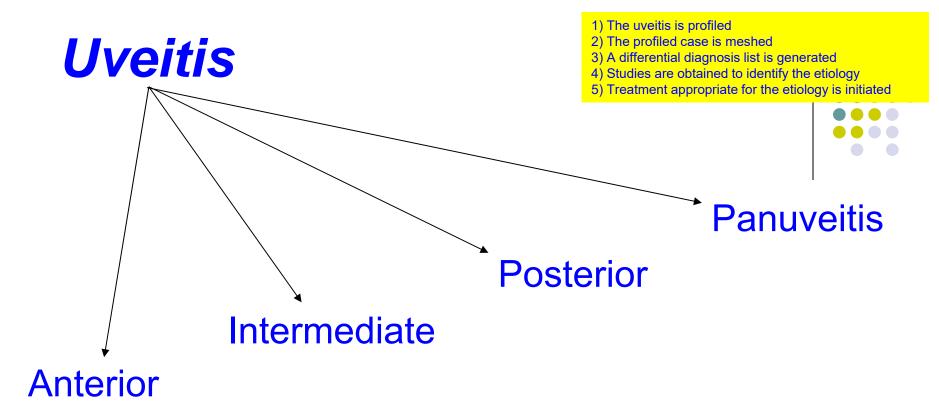
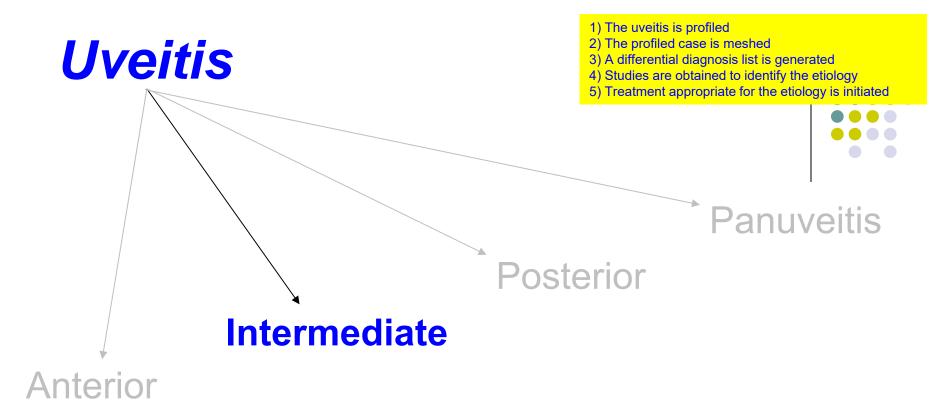
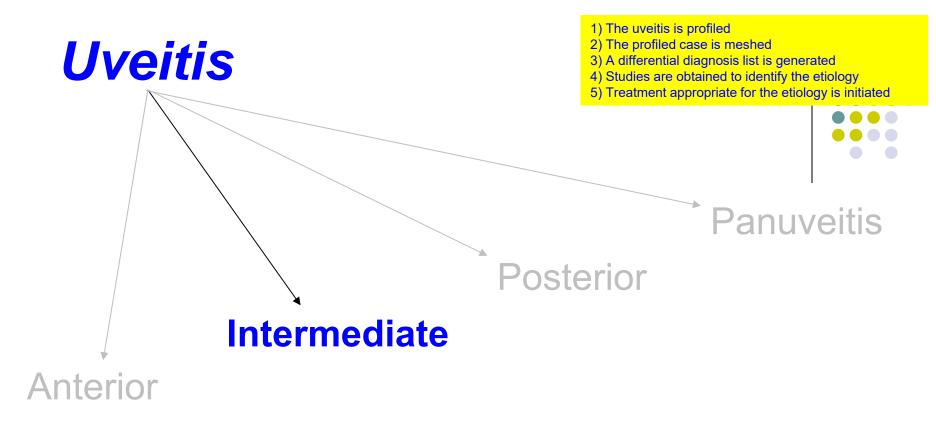


What are the four basic anatomic locations for uveitis?

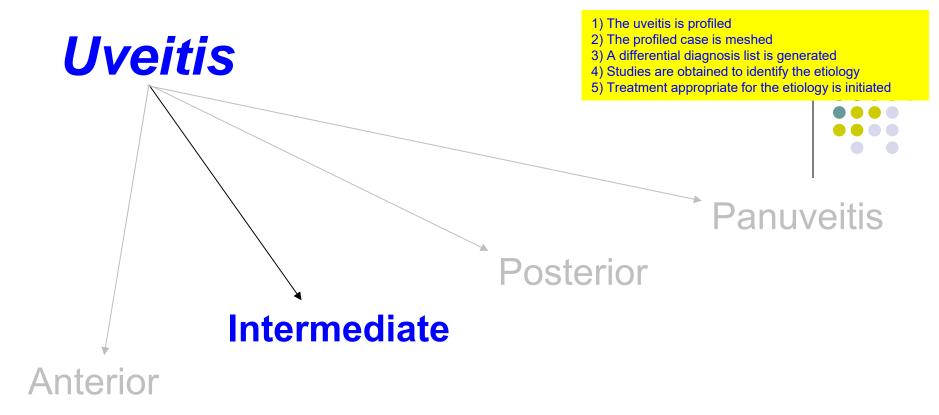


What are the four basic anatomic locations for uveitis?





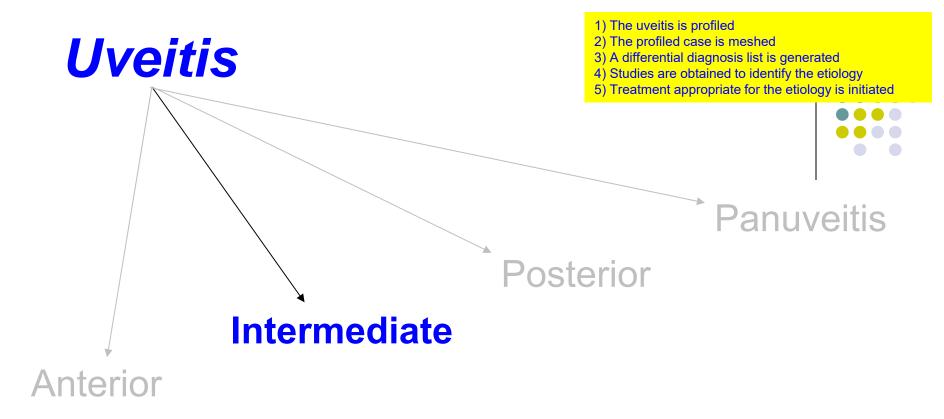
The inflammation is located principally in the anterior vitreous, and usually involves the vitreous base



The inflammation is located principally in the anterior vitreous, and usually involves the vitreous base



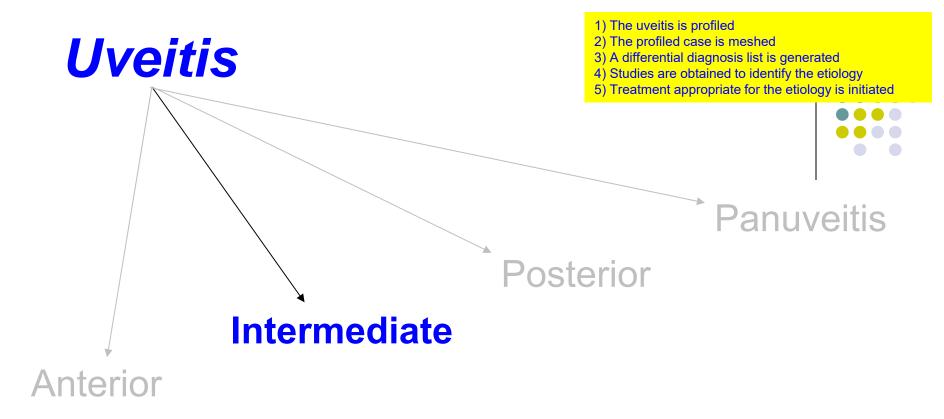
The vitreous base straddles/adheres to what structure?



The inflammation is located principally in the anterior vitreous, and usually involves the vitreous base



The vitreous base straddles/adheres to what structure? The ora serrata

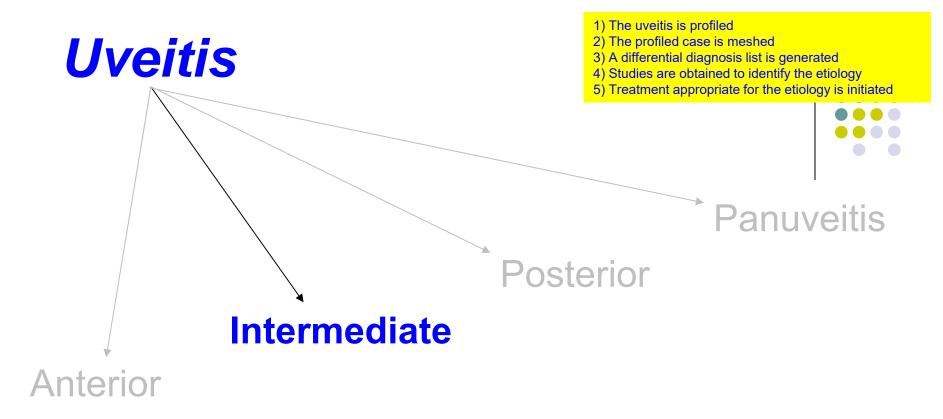


The inflammation is located principally in the anterior vitreous, and usually involves the vitreous base

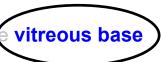


The vitreous base straddles/adheres to what structure? The ora serrata

What is the ora serrata?



The inflammation is located principally in the anterior vitreous, and usually involves the vitreous base



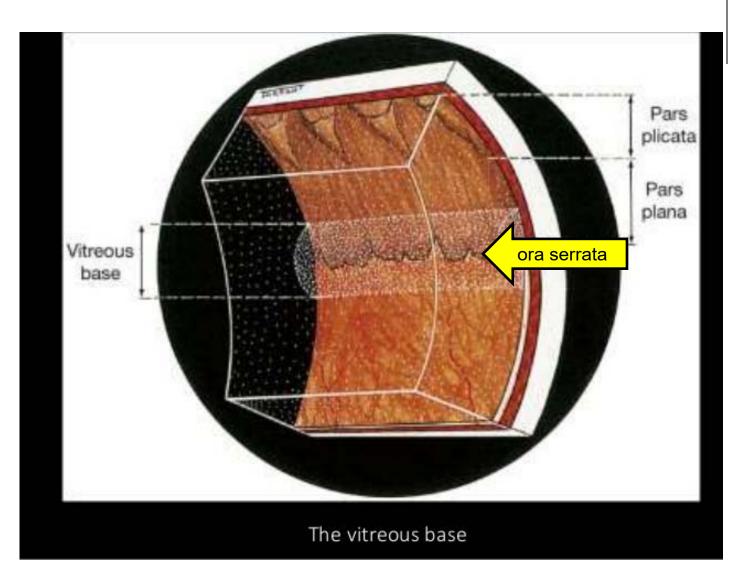
The vitreous base straddles/adheres to what structure?

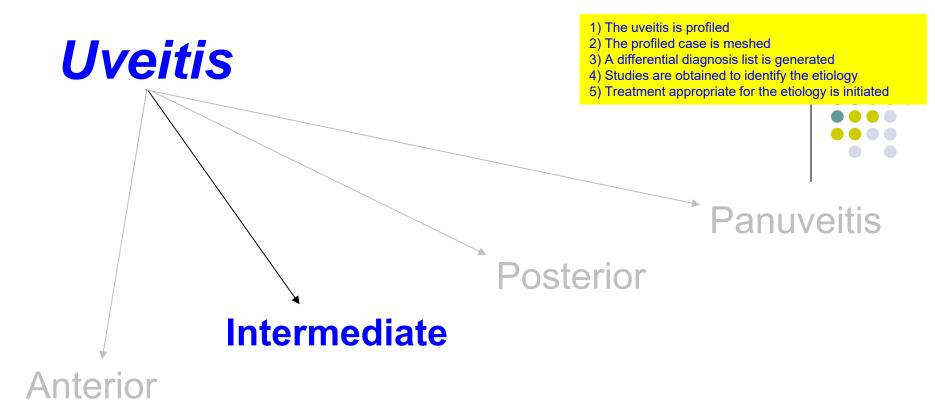
The ora serrata

What is the ora serrata?

It is the zone where the peripheral retina meets the pars plana of the ciliary body

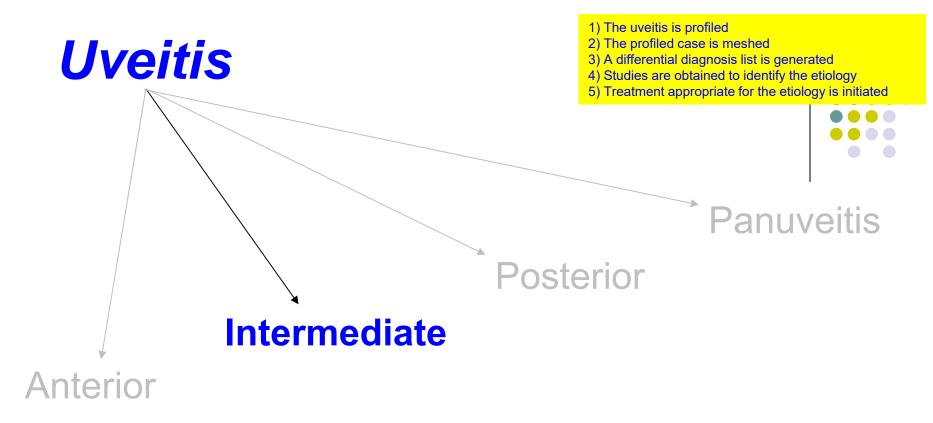






What is the hallmark of intermediate uveitis (IU)? The inflammation is located principally in the anterior vitreous, and usually involves the vitreous base

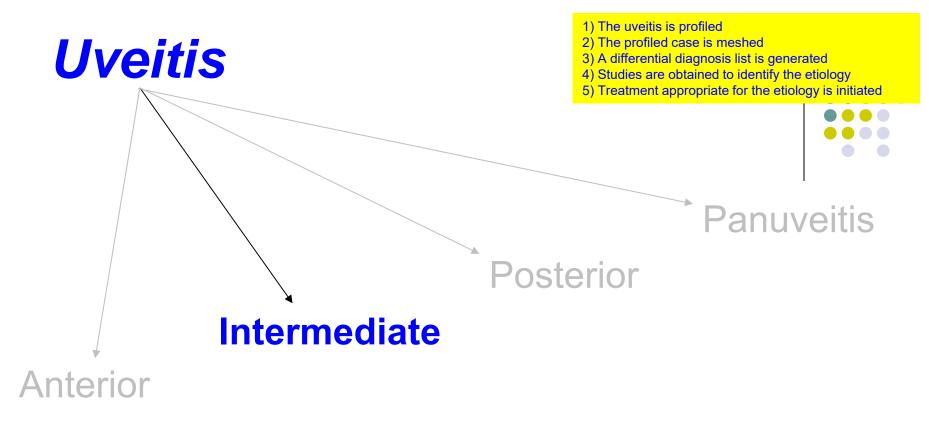
Does IU present with anterior-chamber cell?



The inflammation is located principally in the anterior vitreous, and usually involves the vitreous base

Does IU present with anterior-chamber cell?

Generally no. When present, AC cell is usually mild, and is generally believed to be 'spillover' from the vitreous.

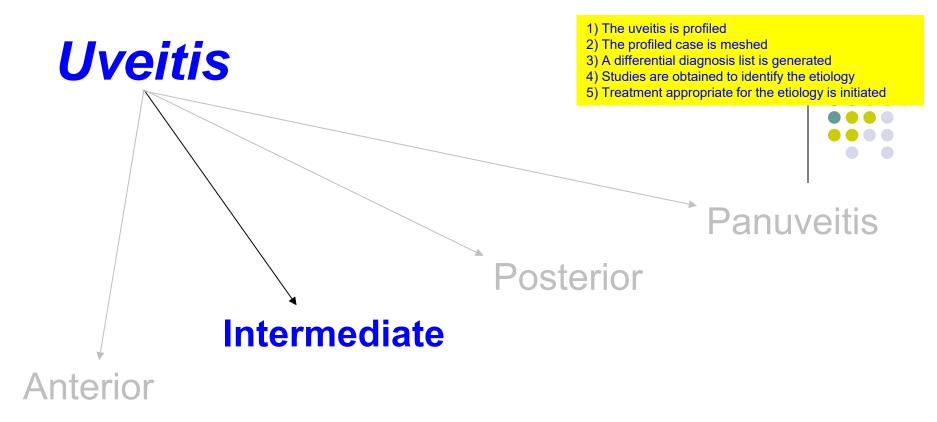


The inflammation is located principally in the anterior vitreous, and usually involves the vitreous base

Does IU present with anterior-chamber cell?

Generally no. When present, AC cell is usually mild, and is generally believed to be 'spillover' from the vitreous.

In this context, what are snowballs and snowbanking?



The inflammation is located principally in the anterior vitreous, and usually involves the vitreous base

Does IU present with anterior-chamber cell?

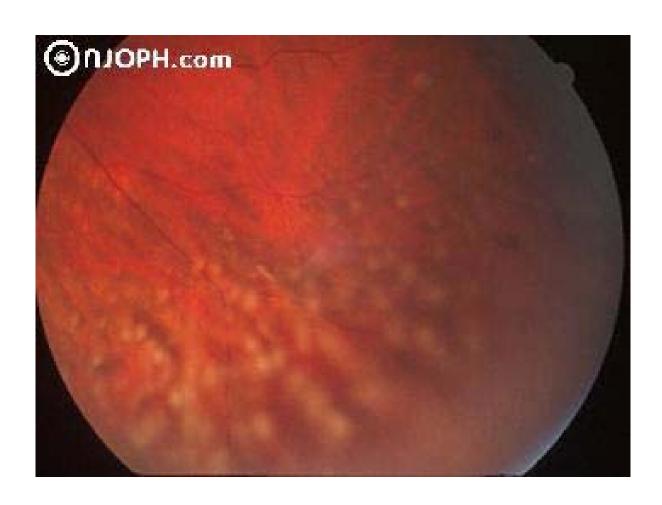
Generally no. When present, AC cell is usually mild, and is generally believed to be 'spillover' from the vitreous.

In this context, what are snowballs and snowbanking?

Snowballs are clumped inflammatory cells in the vitreous

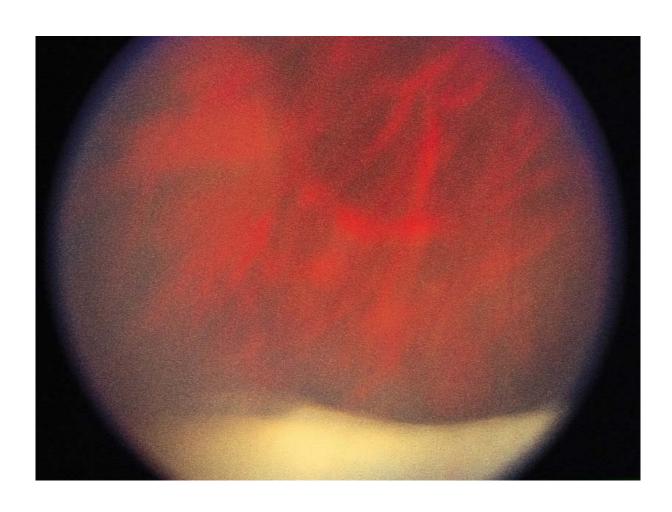
Snowbanking refers to the appearance of large swaths of accumulated inflammatory debris along the inferior pars plana





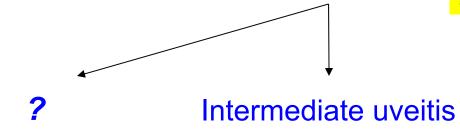
Snowballs in intermediate uveitis





Snowbanking in intermediate uveitis

- 1) The uveitis is profiled
- 2) The profiled case is meshed
- 3) A differential diagnosis list is generated4) Studies are obtained to identify the etiology
- 5) Treatment appropriate for the etiology is initiated



(subtype of intermediate uveitis)



- 1) The uveitis is profiled
- 2) The profiled case is meshed
- 3) A differential diagnosis list is generated
- 4) Studies are obtained to identify the etiology
- 5) Treatment appropriate for the etiology is initiated



Pars planitis

Intermediate uveitis

(subtype of intermediate uveitis)



- 1) The uveitis is profiled
- 2) The profiled case is meshed
- 3) A differential diagnosis list is generated
- 4) Studies are obtained to identify the etiology
- 5) Treatment appropriate for the etiology is initiated

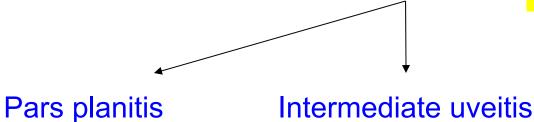


Pars planitis

Intermediate uveitis

When is intermediate uveitis intermediate uveitis, and when is it pars planitis?

- 1) The uveitis is profiled
- 2) The profiled case is meshed
- 3) A differential diagnosis list is generated
- 4) Studies are obtained to identify the etiology
- 5) Treatment appropriate for the etiology is initiated



When is intermediate uveitis intermediate uveitis, and when is it pars planitis? It is considered intermediate uveitis when it is associated with either a local infection or a systemic condition; otherwise it is pars planitis

- 1) The uveitis is profiled
- 2) The profiled case is meshed
- 3) A differential diagnosis list is generated
- 4) Studies are obtained to identify the etiology
- 5) Treatment appropriate for the etiology is initiated



Pars planitis

Intermediate uveitis

At what age(s) is/are pars planitis (PP) most likely to present?

- 1) The uveitis is profiled
- 2) The profiled case is meshed
- 3) A differential diagnosis list is generated
- 4) Studies are obtained to identify the etiology
- 5) Treatment appropriate for the etiology is initiated



• • • • •

Pars planitis

Intermediate uveitis

At what age(s) is/are pars planitis (PP) most likely to present?

The age-of-incidence for PP is bimodal, with one peak in the y.o. range

y.o. range

- 1) The uveitis is profiled
- 2) The profiled case is meshed
- 3) A differential diagnosis list is generated
- 4) Studies are obtained to identify the etiology
- 5) Treatment appropriate for the etiology is initiated





Pars planitis

Intermediate uveitis

At what age(s) is/are pars planitis (PP) most likely to present?

The age-of-incidence for PP is bimodal, with one peak in the 5-15 year-old range, and another in the 20-40 y.o. range

- 1) The uveitis is profiled
- 2) The profiled case is meshed
- 3) A differential diagnosis list is generated
- 4) Studies are obtained to identify the etiology
- 5) Treatment appropriate for the etiology is initiated





Pars planitis

Intermediate uveitis

At what age(s) is/are pars planitis (PP) most likely to present? The age-of-incidence for PP is bimodal, with one peak in the 5-15 year-old range, and another in the 20-40 y.o. range

Is there a gender predilection?

- 1) The uveitis is profiled
- 2) The profiled case is meshed
- 3) A differential diagnosis list is generated
- 4) Studies are obtained to identify the etiology
- 5) Treatment appropriate for the etiology is initiated





Pars planitis

Intermediate uveitis

At what age(s) is/are pars planitis (PP) most likely to present?

The age-of-incidence for PP is bimodal, with one peak in the 5-15 year-old range, and another in the 20-40 y.o. range

Is there a gender predilection?
No

- 1) The uveitis is profiled
- 2) The profiled case is meshed
- 3) A differential diagnosis list is generated
- 4) Studies are obtained to identify the etiology
- 5) Treatment appropriate for the etiology is initiated





Pars planitis

Intermediate uveitis

At what age(s) is/are pars planitis (PP) most likely to present? The age-of-incidence for PP is bimodal, with one peak in the 5-15 year-old range, and another in the 20-40 y.o. range

Is there a gender predilection?
No

What is the typical presenting complaint?

- 1) The uveitis is profiled
- 2) The profiled case is meshed
- 3) A differential diagnosis list is generated
- 4) Studies are obtained to identify the etiology
- 5) Treatment appropriate for the etiology is initiated





Pars planitis

Intermediate uveitis

At what age(s) is/are pars planitis (PP) most likely to present? The age-of-incidence for PP is bimodal, with one peak in the 5-15 year-old range, and another in the 20-40 y.o. range

*Is there a gender predilection?*No

What is the typical presenting complaint?

Children/teens often present in a manner similar to acute anterior uveitis:



and

- 1) The uveitis is profiled
- 2) The profiled case is meshed
- 3) A differential diagnosis list is generated
- 4) Studies are obtained to identify the etiology
- 5) Treatment appropriate for the etiology is initiated





Pars planitis

Intermediate uveitis

At what age(s) is/are pars planitis (PP) most likely to present? The age-of-incidence for PP is bimodal, with one peak in the 5-15 year-old range, and another in the 20-40 y.o. range

Is there a gender predilection?
No

What is the typical presenting complaint? Children/teens often present in a manner similar to acute anterior uveitis: Pain, injection and photophobia.

- 1) The uveitis is profiled
- 2) The profiled case is meshed
- 3) A differential diagnosis list is generated
- 4) Studies are obtained to identify the etiology
- 5) Treatment appropriate for the etiology is initiated





Pars planitis

Intermediate uveitis

At what age(s) is/are pars planitis (PP) most likely to present? The age-of-incidence for PP is bimodal, with one peak in the 5-15 year-old range, and another in the 20-40 y.o. range

Is there a gender predilection?
No

What is the typical presenting complaint?
Children/teens often present in a manner similar to acute anterior uveitis: Pain, injection and photophobia. Adult-onset PP typically presents more insidiously, with a chief complaint of

- 1) The uveitis is profiled
- 2) The profiled case is meshed
- 3) A differential diagnosis list is generated
- 4) Studies are obtained to identify the etiology
- 5) Treatment appropriate for the etiology is initiated





Pars planitis

Intermediate uveitis

At what age(s) is/are pars planitis (PP) most likely to present? The age-of-incidence for PP is bimodal, with one peak in the 5-15 year-old range, and another in the 20-40 y.o. range

Is there a gender predilection?
No

What is the typical presenting complaint?

Children/teens often present in a manner similar to acute anterior uveitis: Pain, injection and photophobia. Adult-onset PP typically presents more insidiously, with a chief complaint of floaters.

- 1) The uveitis is profiled
- 2) The profiled case is meshed
- 3) A differential diagnosis list is generated
- 4) Studies are obtained to identify the etiology
- 5) Treatment appropriate for the etiology is initiated





Pars planitis

Intermediate uveitis

At what age(s) is/are pars planitis (PP) most likely to present? The age-of-incidence for PP is bimodal, with one peak in the 5-15 year-old range, and another in the 20-40 y.o. range

Is there a gender predilection?

What is the typical presenting complaint?
Children/teens often present in a manner similar to acute anterior uveitis: Pain, injection and photophobia. Adult-onset PP typically presents more insidiously, with a chief complaint of floaters.

Does PP tend to present unilaterally, or bilaterally?

- 1) The uveitis is profiled
- 2) The profiled case is meshed
- 3) A differential diagnosis list is generated
- 4) Studies are obtained to identify the etiology
- 5) Treatment appropriate for the etiology is initiated





Pars planitis

Intermediate uveitis

At what age(s) is/are pars planitis (PP) most likely to present? The age-of-incidence for PP is bimodal, with one peak in the 5-15 year-old range, and another in the 20-40 y.o. range

Is there a gender predilection?

What is the typical presenting complaint?

Children/teens often present in a manner similar to acute anterior uveitis: Pain, injection and photophobia. Adult-onset PP typically presents more insidiously, with a chief complaint of floaters.

Does PP tend to present unilaterally, or bilaterally?

It is bilateral in 6 of cases (but it is often quite asymmetric in severity)

- 1) The uveitis is profiled
- 2) The profiled case is meshed
- 3) A differential diagnosis list is generated
- 4) Studies are obtained to identify the etiology
- 5) Treatment appropriate for the etiology is initiated





Pars planitis

Intermediate uveitis

At what age(s) is/are pars planitis (PP) most likely to present? The age-of-incidence for PP is bimodal, with one peak in the 5-15 year-old range, and another in the 20-40 y.o. range

Is there a gender predilection?

What is the typical presenting complaint?

Children/teens often present in a manner similar to acute anterior uveitis: Pain, injection and photophobia. Adult-onset PP typically presents more insidiously, with a chief complaint of floaters.

Does PP tend to present unilaterally, or bilaterally?

It is bilateral in 80% of cases (but it is often quite asymmetric in severity)

- 1) The uveitis is profiled
- 2) The profiled case is meshed
- 3) A differential diagnosis list is generated
- 4) Studies are obtained to identify the etiology
- 5) Treatment appropriate for the etiology is initiated





Pars planitis

Intermediate uveitis

At what age(s) is/are pars planitis (PP) most likely to present? The age-of-incidence for PP is bimodal, with one peak in the 5-15 year-old range, and another in the 20-40 y.o. range

Is there a gender predilection?

What is the typical presenting complaint?

Children/teens often present in a manner similar to acute anterior uveitis: Pain, injection and photophobia. Adult-onset PP typically presents more insidiously, with a chief complaint of floaters.

Does PP tend to present unilaterally, or bilaterally?

It is bilateral in 80% of cases (but it is often quite asymmetric in severity)

Is PP a benign condition?

- 1) The uveitis is profiled
- 2) The profiled case is meshed
- 3) A differential diagnosis list is generated
- 4) Studies are obtained to identify the etiology
- 5) Treatment appropriate for the etiology is initiated





Pars planitis

Intermediate uveitis

At what age(s) is/are pars planitis (PP) most likely to present? The age-of-incidence for PP is bimodal, with one peak in the 5-15 year-old range, and another in the 20-40 y.o. range

Is there a gender predilection?

What is the typical presenting complaint?

Children/teens often present in a manner similar to acute anterior uveitis: Pain, injection and photophobia. Adult-onset PP typically presents more insidiously, with a chief complaint of floaters.

Does PP tend to present unilaterally, or bilaterally?

It is bilateral in 80% of cases (but it is often quite asymmetric in severity)

Is PP a benign condition?
Unfortunately, no. There are two significant sequelae to worry about:

__

- 1) The uveitis is profiled
- 2) The profiled case is meshed
- 3) A differential diagnosis list is generated
- 4) Studies are obtained to identify the etiology
- 5) Treatment appropriate for the etiology is initiated





Pars planitis

Intermediate uveitis

At what age(s) is/are pars planitis (PP) most likely to present? The age-of-incidence for PP is bimodal, with one peak in the 5-15 year-old range, and another in the 20-40 y.o. range

Is there a gender predilection?

What is the typical presenting complaint?

Children/teens often present in a manner similar to acute anterior uveitis: Pain, injection and photophobia. Adult-onset PP typically presents more insidiously, with a chief complaint of floaters.

Does PP tend to present unilaterally, or bilaterally?

It is bilateral in 80% of cases (but it is often quite asymmetric in severity)

Is PP a benign condition?

Unfortunately, no. There are two significant sequelae to worry about:

- --CME
- -- Neovascularization of the far-peripheral retina

- 1) The uveitis is profiled
- 2) The profiled case is meshed
- 3) A differential diagnosis list is generated
- 4) Studies are obtained to identify the etiology
- 5) Treatment appropriate for the etiology is initiated





Pars planitis

Intermediate uveitis

At what age(s) is/are pars planitis (PP) most likely to present? The age-of-incidence for PP is bimodal, with one peak in the 5-15 year-old range, and another in the 20-40 y.o. range

Is there a gender predilection?

What is the typical presenting complaint?

Children/teens often present in a manner similar to acute anterior uveitis: Pain, injection and photophobia. Adult-onset PP typically presents more insidiously, with a chief complaint of floaters.

Does PP tend to present unilaterally, or bilaterally?

It is bilateral in 80% of cases (but it is often quite asymmetric in severity)

Is PP a benign condition?

Unfortunately, no. There are two significant sequelae to worry about:

- --CME is common v uncommon
- --Neovascularization of the far-peripheral retina

- 1) The uveitis is profiled
- 2) The profiled case is meshed
- 3) A differential diagnosis list is generated
- 4) Studies are obtained to identify the etiology
- 5) Treatment appropriate for the etiology is initiated





Pars planitis

Intermediate uveitis

At what age(s) is/are pars planitis (PP) most likely to present? The age-of-incidence for PP is bimodal, with one peak in the 5-15 year-old range, and another in the 20-40 y.o. range

Is there a gender predilection?

What is the typical presenting complaint?

Children/teens often present in a manner similar to acute anterior uveitis: Pain, injection and photophobia. Adult-onset PP typically presents more insidiously, with a chief complaint of floaters.

Does PP tend to present unilaterally, or bilaterally?

It is bilateral in 80% of cases (but it is often quite asymmetric in severity)

Is PP a benign condition?

- -- CME is common
- --Neovascularization of the far-peripheral retina

- 1) The uveitis is profiled
- 2) The profiled case is meshed
- 3) A differential diagnosis list is generated
- 4) Studies are obtained to identify the etiology
- 5) Treatment appropriate for the etiology is initiated





Pars planitis

Intermediate uveitis

At what age(s) is/are pars planitis (PP) most likely to present? The age-of-incidence for PP is bimodal, with one peak in the 5-15 year-old range, and another in the 20-40 y.o. range

Is there a gender predilection?

What is the typical presenting complaint?

Children/teens often present in a manner similar to acute anterior uveitis: Pain, injection and photophobia. Adult-onset PP typically presents more insidiously, with a chief complaint of floaters.

Does PP tend to present unilaterally, or bilaterally?

It is bilateral in 80% of cases (but it is often quite asymmetric in severity)

Is PP a benign condition?

- --CME is common, and leads to permanent vision loss in about 6 of cases
- --Neovascularization of the far-peripheral retina

- 1) The uveitis is profiled
- 2) The profiled case is meshed
- 3) A differential diagnosis list is generated
- 4) Studies are obtained to identify the etiology
- 5) Treatment appropriate for the etiology is initiated





Pars planitis

Intermediate uveitis

At what age(s) is/are pars planitis (PP) most likely to present? The age-of-incidence for PP is bimodal, with one peak in the 5-15 year-old range, and another in the 20-40 y.o. range

Is there a gender predilection?

What is the typical presenting complaint?

Children/teens often present in a manner similar to acute anterior uveitis: Pain, injection and photophobia. Adult-onset PP typically presents more insidiously, with a chief complaint of floaters.

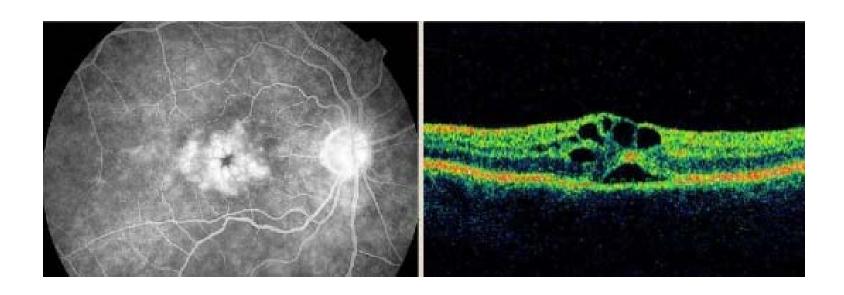
Does PP tend to present unilaterally, or bilaterally?

It is bilateral in 80% of cases (but it is often quite asymmetric in severity)

Is PP a benign condition?

- -- CME is common, and leads to permanent vision loss in about 10% of cases
- --Neovascularization of the far-peripheral retina





CME in intermediate uveitis

- 1) The uveitis is profiled
- 2) The profiled case is meshed
- 3) A differential diagnosis list is generated
- 4) Studies are obtained to identify the etiology
- 5) Treatment appropriate for the etiology is initiated





Pars planitis

Intermediate uveitis

At what age(s) is/are pars planitis (PP) most likely to present? The age-of-incidence for PP is bimodal, with one peak in the 5-15 year-old range, and another in the 20-40 y.o. range

Is there a gender predilection?

What is the typical presenting complaint?

Children/teens often present in a manner similar to acute anterior uveitis: Pain, injection and photophobia. Adult-onset PP typically presents more insidiously, with a chief complaint of floaters.

Does PP tend to present unilaterally, or bilaterally?

It is bilateral in 80% of cases (but it is often quite asymmetric in severity)

Is PP a benign condition?

Unfortunately, no. There are two significant sequelae to worry about:

- -- CME is common, and leads to permanent vision loss in about 10% of cases
- --Neovascularization of the far-peripheral retina may occur with all its attendant problems:

two words ; two different words ; word-abb and/or diff word, same abb

- 1) The uveitis is profiled
- 2) The profiled case is meshed
- 3) A differential diagnosis list is generated
- 4) Studies are obtained to identify the etiology
- 5) Treatment appropriate for the etiology is initiated





Pars planitis

Intermediate uveitis

At what age(s) is/are pars planitis (PP) most likely to present? The age-of-incidence for PP is bimodal, with one peak in the 5-15 year-old range, and another in the 20-40 y.o. range

Is there a gender predilection?

What is the typical presenting complaint?

Children/teens often present in a manner similar to acute anterior uveitis: Pain, injection and photophobia. Adult-onset PP typically presents more insidiously, with a chief complaint of floaters.

Does PP tend to present unilaterally, or bilaterally?

It is bilateral in 80% of cases (but it is often quite asymmetric in severity)

Is PP a benign condition?

- --CME is common, and leads to permanent vision loss in about 10% of cases
- --Neovascularization of the far-peripheral retina may occur with all its attendant problems: vitreous hemorrhage; retinal traction; tractional RD and/or rhegmatogenous RD

- 1) The uveitis is profiled
- 2) The profiled case is meshed
- 3) A differential diagnosis list is generated
- 4) Studies are obtained to identify the etiology
- 5) Treatment appropriate for the etiology is initiated





Pars planitis

Intermediate uveitis

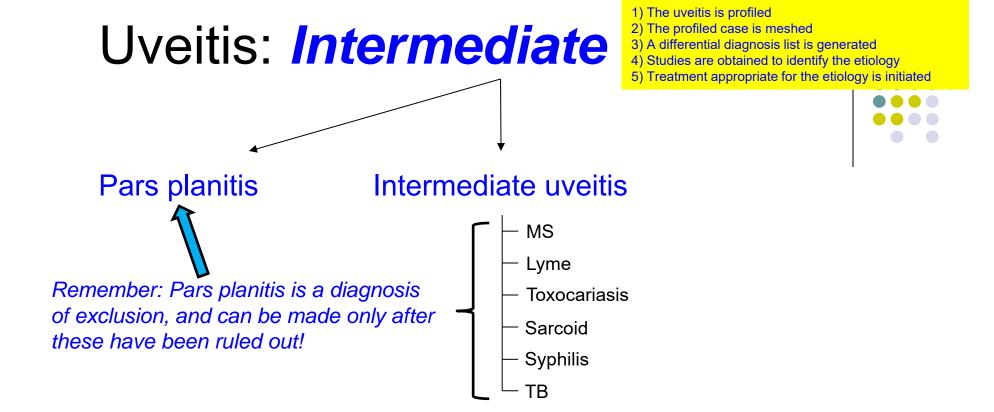
- ? - ? - ? - ?

When is intermediate uveitis intermediate uveitis, and when is it pars planitis? It is considered intermediate uveitis when it is associated with either a local infection or a systemic condition

With what infections/systemic conditions is intermediate uveitis associated?

When is intermediate uveitis intermediate uveitis, and when is it pars planitis? It is considered intermediate uveitis when it is associated with either a local infection or a systemic condition

With what infections/systemic conditions is intermediate uveitis associated?



When is intermediate uveitis intermediate uveitis, and when is it pars planitis? It is considered intermediate uveitis when it is associated with either a local infection or a systemic condition

With what infections/systemic conditions is intermediate uveitis associated?

- 1) The uveitis is profiled
- 2) The profiled case is meshed
- 3) A differential diagnosis list is generated
- 4) Studies are obtained to identify the etiology
- 5) Treatment appropriate for the etiology is initiated



• • • • •

Pars planitis

Intermediate uveitis

How is pars planitis/intermediate uveitis (PP/IU) managed?

- 1) The uveitis is profiled
- 2) The profiled case is meshed
- 3) A differential diagnosis list is generated
- 4) Studies are obtained to identify the etiology
- 5) Treatment appropriate for the etiology is initiated



Pars planitis

Intermediate uveitis

How is pars planitis/intermediate uveitis (PP/IU) managed?

If an etiology is identified (ie, if it is IU), treatment specific to that etiology should be pursued

- 1) The uveitis is profiled
- 2) The profiled case is meshed
- 3) A differential diagnosis list is generated
- 4) Studies are obtained to identify the etiology
- 5) Treatment appropriate for the etiology is initiated





Pars planitis

Intermediate uveitis

How is pars planitis/intermediate uveitis (PP/IU) managed? If an etiology is identified (ie, if it is IU), treatment specific to that etiology should be pursued

Assume testing is noncontributory. How should PP be managed?

- 1) The uveitis is profiled
- 2) The profiled case is meshed
- 3) A differential diagnosis list is generated
- 4) Studies are obtained to identify the etiology
- 5) Treatment appropriate for the etiology is initiated





Pars planitis

Intermediate uveitis

How is pars planitis/intermediate uveitis (PP/IU) managed? If an etiology is identified (ie, if it is IU), treatment specific to that etiology should be pursued

Assume testing is noncontributory. How should PP be managed? If it is mild, and not causing significant morbidity, it can simply be monitored

- 1) The uveitis is profiled
- 2) The profiled case is meshed
- 3) A differential diagnosis list is generated
- 4) Studies are obtained to identify the etiology
- 5) Treatment appropriate for the etiology is initiated





Pars planitis

Intermediate uveitis

How is pars planitis/intermediate uveitis (PP/IU) managed? If an etiology is identified (ie, if it is IU), treatment specific to that etiology should be pursued

Assume testing is noncontributory. How should PP be managed? If it is mild, and not causing significant morbidity, it can simply be monitored

Under what circumstances should treatment be initiated?

- 1) The uveitis is profiled
- 2) The profiled case is meshed
- 3) A differential diagnosis list is generated
- 4) Studies are obtained to identify the etiology
- 5) Treatment appropriate for the etiology is initiated



Pars planitis

Intermediate uveitis

How is pars planitis/intermediate uveitis (PP/IU) managed?

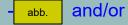
If an etiology is identified (ie, if it is IU), treatment specific to that etiology should be pursued

Assume testing is noncontributory. How should PP be managed? If it is mild, and not causing significant morbidity, it can simply be monitored

Under what circumstances should treatment be initiated?

Treatment should initiated if:

-- The pt's vision is affected; or



two words

develops

- 1) The uveitis is profiled
- 2) The profiled case is meshed
- 3) A differential diagnosis list is generated
- 4) Studies are obtained to identify the etiology
- 5) Treatment appropriate for the etiology is initiated





Pars planitis

Intermediate uveitis

How is pars planitis/intermediate uveitis (PP/IU) managed?

If an etiology is identified (ie, if it is IU), treatment specific to that etiology should be pursued

Assume testing is noncontributory. How should PP be managed? If it is mild, and not causing significant morbidity, it can simply be monitored

Under what circumstances should treatment be initiated?

Treatment should initiated if:

- -- The pt's vision is affected; or
- -- CME and/or retinal vasculitis develops

- 1) The uveitis is profiled
- 2) The profiled case is meshed
- 3) A differential diagnosis list is generated
- 4) Studies are obtained to identify the etiology
- 5) Treatment appropriate for the etiology is initiated





Pars planitis

Intermediate uveitis

How is pars planitis/intermediate uveitis (PP/IU) managed?

If an etiology is identified (ie, if it is IU), treatment specific to that etiology should be pursued

Assume testing is noncontributory. How should PP be managed? If it is mild, and not causing significant morbidity, it can simply be monitored

Under what circumstances should treatment be initiated?

Treatment should initiated if:

- -- The pt's vision is affected; or
- -- CME and/or retinal vasculitis develops

What sort of treatment plan should be initiated?

- 1) The uveitis is profiled
- 2) The profiled case is meshed
- 3) A differential diagnosis list is generated
- 4) Studies are obtained to identify the etiology
- 5) Treatment appropriate for the etiology is initiated





Pars planitis

Intermediate uveitis

How is pars planitis/intermediate uveitis (PP/IU) managed?

If an etiology is identified (ie, if it is IU), treatment specific to that etiology should be pursued

Assume testing is noncontributory. How should PP be managed? If it is mild, and not causing significant morbidity, it can simply be monitored

Under what circumstances should treatment be initiated?

Treatment should initiated if:

- -- The pt's vision is affected; or
- -- CME and/or retinal vasculitis develops

What sort of treatment plan should be initiated?

- 1) The uveitis is profiled
- 2) The profiled case is meshed
- 3) A differential diagnosis list is generated
- 4) Studies are obtained to identify the etiology
- 5) Treatment appropriate for the etiology is initiated





Pars planitis

Intermediate uveitis

How is pars planitis/intermediate uveitis (PP/IU) managed?

If an etiology is identified (ie, if it is IU), treatment specific to that etiology should be pursued

Assume testing is noncontributory. How should PP be managed? If it is mild, and not causing significant morbidity, it can simply be monitored

Under what circumstances should treatment be initiated?

Treatment should initiated if:

- -- The pt's vision is affected; or
- -- CME and/or retinal vasculitis develops

What sort of treatment plan should be initiated?

- 1)?
- 2)
- 3)
- 4)

- 1) The uveitis is profiled
- 2) The profiled case is meshed
- 3) A differential diagnosis list is generated
- 4) Studies are obtained to identify the etiology
- 5) Treatment appropriate for the etiology is initiated





Pars planitis

Intermediate uveitis

How is pars planitis/intermediate uveitis (PP/IU) managed? If an etiology is identified (ie, if it is IU), treatment specific to that etiology should be pursued

Assume testing is noncontributory. How should PP be managed? If it is mild, and not causing significant morbidity, it can simply be monitored

Under what circumstances should treatment be initiated?

Treatment should initiated if:

- -- The pt's vision is affected; or
- -- CME and/or retinal vasculitis develops

What sort of treatment plan should be initiated?

- 1) Steroid therapy
- 2) ?
- 3)
- 4)

- 1) The uveitis is profiled
- 2) The profiled case is meshed
- 3) A differential diagnosis list is generated
- 4) Studies are obtained to identify the etiology
- 5) Treatment appropriate for the etiology is initiated





Pars planitis

Intermediate uveitis

How is pars planitis/intermediate uveitis (PP/IU) managed?

If an etiology is identified (ie, if it is IU), treatment specific to that etiology should be pursued

Assume testing is noncontributory. How should PP be managed? If it is mild, and not causing significant morbidity, it can simply be monitored

Under what circumstances should treatment be initiated?

Treatment should initiated if:

- -- The pt's vision is affected; or
- -- CME and/or retinal vasculitis develops

What sort of treatment plan should be initiated?

- 1) Steroid therapy
- 2) Peripheral retina ablation
- 3)?
- 4)

- 1) The uveitis is profiled
- 2) The profiled case is meshed
- 3) A differential diagnosis list is generated
- 4) Studies are obtained to identify the etiology
- 5) Treatment appropriate for the etiology is initiated



Pars planitis

Intermediate uveitis

How is pars planitis/intermediate uveitis (PP/IU) managed?

If an etiology is identified (ie, if it is IU), treatment specific to that etiology should be pursued

Assume testing is noncontributory. How should PP be managed? If it is mild, and not causing significant morbidity, it can simply be monitored

Under what circumstances should treatment be initiated?

Treatment should initiated if:

- -- The pt's vision is affected; or
- -- CME and/or retinal vasculitis develops

What sort of treatment plan should be initiated?

- 1) Steroid therapy
- 2) Peripheral retina ablation
- 3) Vitrectomy
- 4)?

- 1) The uveitis is profiled
- 2) The profiled case is meshed
- 3) A differential diagnosis list is generated
- 4) Studies are obtained to identify the etiology
- 5) Treatment appropriate for the etiology is initiated





Pars planitis

Intermediate uveitis

How is pars planitis/intermediate uveitis (PP/IU) managed?

If an etiology is identified (ie, if it is IU), treatment specific to that etiology should be pursued

Assume testing is noncontributory. How should PP be managed? If it is mild, and not causing significant morbidity, it can simply be monitored

Under what circumstances should treatment be initiated?

Treatment should initiated if:

- -- The pt's vision is affected; or
- -- CME and/or retinal vasculitis develops

What sort of treatment plan should be initiated?

- 1) Steroid therapy
- 2) Peripheral retina ablation
- 3) Vitrectomy
- 4) Immunomodulatory therapy

- 1) The uveitis is profiled
- 2) The profiled case is meshed
- 3) A differential diagnosis list is generated
- 4) Studies are obtained to identify the etiology
- 5) Treatment appropriate for the etiology is initiated



Pars planitis

Intermediate uveitis

How is pars planitis/intermediate uveitis (PP/IU) managed? If an etiology is identified (ie, if it is IU), treatment specific to that etiology should be pursued

Assume testing is noncontributory. How should PP be managed? If it is mild, and not causing significant morbidity, it can simply be monitored

Under what circumstances should treatment be initiated?

Treatment should initiated if:

- --The pt's vision is affected; or
- -- CME and/or retinal vasculitis develops

What sort of treatment plan should be initiated?

A four-step approach should be employed:

- 1) Steroid therapy
- 2) Peripheral retina ablation
- 3) Vitrectomy
- 4) Immunomodulatory therapy

Some experts would reverse the order of these two

- 1) The uveitis is profiled
- 2) The profiled case is meshed
- 3) A differential diagnosis list is generated
- 4) Studies are obtained to identify the etiology
- 5) Treatment appropriate for the etiology is initiated





Pars planitis

Intermediate uveitis

How is pars planitis/intermediate uveitis (PP/IU) managed? If an etiology is identified (ie, if it is IU), treatment specific to that etiology should be pursued

What is the preferred route of steroid administration?

vvnat son or treatment plan should be initiated:

A four step approach should be employed:

- 2) Peripheral retina ablation
- 3) Vitrectomy
- 4) Immunomodulatory therapy

- 1) The uveitis is profiled
- 2) The profiled case is meshed
- 3) A differential diagnosis list is generated
- 4) Studies are obtained to identify the etiology
- 5) Treatment appropriate for the etiology is initiated





Pars planitis

Intermediate uveitis

How is pars planitis/intermediate uveitis (PP/IU) managed? If an etiology is identified (ie, if it is IU), treatment specific to that etiology should be pursued

What is the preferred route of steroid administration? Periocular depot injection

vvnat son or treatment plan should be initiated?

A <u>four step approac</u>h should be employed:

- 2) Peripheral retina ablation
- 3) Vitrectomy
- 4) Immunomodulatory therapy

- 1) The uveitis is profiled
- 2) The profiled case is meshed
- 3) A differential diagnosis list is generated
- 4) Studies are obtained to identify the etiology
- 5) Treatment appropriate for the etiology is initiated





Pars planitis

Intermediate uveitis

How is pars planitis/intermediate uveitis (PP/IU) managed? If an etiology is identified (ie, if it is IU), treatment specific to that etiology should be pursued

What is the preferred route of steroid administration? Periocular depot injection

What about intravitreal injection?

vvnat sort or treatment plan should be initiated?

A <u>four step approac</u>h should be employed:

- 2) Peripheral retina ablation
- 3) Vitrectomy
- 4) Immunomodulatory therapy

- 1) The uveitis is profiled
- 2) The profiled case is meshed
- 3) A differential diagnosis list is generated
- 4) Studies are obtained to identify the etiology
- 5) Treatment appropriate for the etiology is initiated





Pars planitis

Intermediate uveitis

How is pars planitis/intermediate uveitis (PP/IU) managed?

If an etiology is identified (ie, if it is IU), treatment specific to that etiology should be pursued

What is the preferred route of steroid administration? Periocular depot injection

What about intravitreal injection?

This can be attempted in severe/refractory cases, but care must be taken to not pierce the pars plana at an inflamed location

vvnat sort or treatment plan should be initiated?

A four step approach should be employed:

- 2) Peripheral retina ablation
- 3) Vitrectomy
- 4) Immunomodulatory therapy

- 1) The uveitis is profiled
- 2) The profiled case is meshed
- 3) A differential diagnosis list is generated
- 4) Studies are obtained to identify the etiology
- 5) Treatment appropriate for the etiology is initiated





Pars planitis

Intermediate uveitis

How is pars planitis/intermediate uveitis (PP/IU) managed?

If an etiology is identified (ie, if it is IU), treatment specific to that etiology should be pursued

What is the preferred route of steroid administration? Periocular depot injection

What about intravitreal injection?

This can be attempted in severe/refractory cases, but care must be taken to not pierce the pars plana at an inflamed location

What about systemic steroids?

vvriat sort or treatment plan should be initiated?

A <u>four etep approac</u>h should be employed:

- 2) Peripheral retina ablation
- 3) Vitrectomy
- 4) Immunomodulatory therapy

- 1) The uveitis is profiled
- 2) The profiled case is meshed
- 3) A differential diagnosis list is generated
- 4) Studies are obtained to identify the etiology
- 5) Treatment appropriate for the etiology is initiated





Pars planitis

Intermediate uveitis

How is pars planitis/intermediate uveitis (PP/IU) managed?
If an etiology is identified (ie, if it is IU), treatment specific to that etiology should be pursued

What is the preferred route of steroid administration?

Periocular depot injection

What about intravitreal injection?

This can be attempted in severe/refractory cases, but care must be taken to not pierce the pars plana at an inflamed location

What about systemic steroids?

These can also be considered for severe/refractory disease, especially in

cases

vvnat sort or treatment plan snould be initiated?

A four step approach should be employed:

- 2) Peripheral retina ablation
- 3) Vitrectomy
- 4) Immunomodulatory therapy

- 1) The uveitis is profiled
- 2) The profiled case is meshed
- 3) A differential diagnosis list is generated
- 4) Studies are obtained to identify the etiology
- 5) Treatment appropriate for the etiology is initiated





Pars planitis

Intermediate uveitis

How is pars planitis/intermediate uveitis (PP/IU) managed? If an etiology is identified (ie, if it is IU), treatment specific to that etiology should be pursued

What is the preferred route of steroid administration? Periocular depot injection

What about intravitreal injection?

This can be attempted in severe/refractory cases, but care must be taken to not pierce the pars plana at an inflamed location

What about systemic steroids?

These can also be considered for severe/refractory disease, especially in bilateral cases

vvnat son or treatment plan should be initiated?

A <u>four etep approac</u>h should be employed:

- 2) Peripheral retina ablation
- 3) Vitrectomy
- 4) Immunomodulatory therapy

- 1) The uveitis is profiled
- 2) The profiled case is meshed
- 3) A differential diagnosis list is generated
- 4) Studies are obtained to identify the etiology
- 5) Treatment appropriate for the etiology is initiated



Pars planitis

Intermediate uveitis

How is pars planitis/intermediate uveitis (PP/IU) managed?

When should peripheral retinal ablation be pursued?

ursued

- 1) Steroid therapy
- 2) Peripheral retina ablation
- 3) Vitrectomy
- 4) Immunomodulatory therapy

- 1) The uveitis is profiled
- 2) The profiled case is meshed
- 3) A differential diagnosis list is generated
- 4) Studies are obtained to identify the etiology
- 5) Treatment appropriate for the etiology is initiated



Pars planitis

Intermediate uveitis

How is pars planitis/intermediate uveitis (PP/IU) managed?

When should peripheral retinal ablation be pursued? If the pt fails steroid therapy

ursued

- 1) Steroid therapy
- 2) Peripheral retina ablation
- 3) Vitrectomy
- 4) Immunomodulatory therapy

- 1) The uveitis is profiled
- 2) The profiled case is meshed
- 3) A differential diagnosis list is generated
- 4) Studies are obtained to identify the etiology
- 5) Treatment appropriate for the etiology is initiated



Pars planitis

Intermediate uveitis

How is pars planitis/intermediate uveitis (PP/IU) managed?

When should peripheral retinal ablation be pursued? If the pt fails steroid therapy

What modality(ies) can be employed?

- 1) Steroid therapy
- 2) Peripheral retina ablation
- 3) Vitrectomy
- 4) Immunomodulatory therapy

- 1) The uveitis is profiled
- 2) The profiled case is meshed
- 3) A differential diagnosis list is generated
- 4) Studies are obtained to identify the etiology
- 5) Treatment appropriate for the etiology is initiated



Pars planitis

Intermediate uveitis

How is pars planitis/intermediate uveitis (PP/IU) managed?

When should peripheral retinal ablation be pursued? If the pt fails steroid therapy

What modality(ies) can be employed? Cryoablation, or laser photocoagulation

- 1) Steroid therapy
- 2) Peripheral retina ablation
- 3) Vitrectomy
- 4) Immunomodulatory therapy

irsued

- 1) The uveitis is profiled
- 2) The profiled case is meshed
- 3) A differential diagnosis list is generated
- 4) Studies are obtained to identify the etiology
- 5) Treatment appropriate for the etiology is initiated



Pars planitis

Intermediate uveitis

How is pars planitis/intermediate uveitis (PP/IU) managed?

When should peripheral retinal ablation be pursued? If the pt fails steroid therapy

What modality(ies) can be employed? Cryoablation, or laser photocoagulation

Which specific area should be treated?

- 1) Steroid therapy
- 2) Peripheral retina ablation
- 3) Vitrectomy
- 4) Immunomodulatory therapy

ursued

- 1) The uveitis is profiled
- 2) The profiled case is meshed
- 3) A differential diagnosis list is generated
- 4) Studies are obtained to identify the etiology
- 5) Treatment appropriate for the etiology is initiated



Pars planitis

Intermediate uveitis

How is pars planitis/intermediate uveitis (PP/IU) managed?

When should peripheral retinal ablation be pursued? If the pt fails steroid therapy

What modality(ies) can be employed? Cryoablation, or laser photocoagulation

Which specific area should be treated? If cryoablation is used, it should be applied...

- 1) Steroid therapy
- 2) Peripheral retina ablation
- 3) Vitrectomy
- 4) Immunomodulatory therapy

ursued

- 1) The uveitis is profiled
- 2) The profiled case is meshed
- 3) A differential diagnosis list is generated
- 4) Studies are obtained to identify the etiology
- 5) Treatment appropriate for the etiology is initiated



Pars planitis

Intermediate uveitis

How is pars planitis/intermediate uveitis (PP/IU) managed?

When should peripheral retinal ablation be pursued? If the pt fails steroid therapy

What modality(ies) can be employed? Cryoablation, or laser photocoagulation

If cryoablation is used, it should be applied...directly to the sclera adjacent to where snowbanking is present

Which specific area should be treated?

- 2) Peripheral retina ablation
- 4) Immunomodulatory therapy

- 1) The uveitis is profiled
- 2) The profiled case is meshed
- 3) A differential diagnosis list is generated
- 4) Studies are obtained to identify the etiology
- 5) Treatment appropriate for the etiology is initiated



Pars planitis

Intermediate uveitis

How is pars planitis/intermediate uveitis (PP/IU) managed?

When should peripheral retinal ablation be pursued? If the pt fails steroid therapy

What modality(ies) can be employed? Cryoablation, or laser photocoagulation

Which specific area should be treated?

If cryoablation is used, it should be applied...directly to the sclera adjacent to where snowbanking is present

If laser photocoagulation is used, it should be applied...

- 1) Steroid therapy
- 2) Peripheral retina ablation
- 3) Vitrectomy
- 4) Immunomodulatory therapy

- 1) The uveitis is profiled
- 2) The profiled case is meshed
- 3) A differential diagnosis list is generated
- 4) Studies are obtained to identify the etiology
- 5) Treatment appropriate for the etiology is initiated



Pars planitis

Intermediate uveitis

How is pars planitis/intermediate uveitis (PP/IU) managed?

When should peripheral retinal ablation be pursued? If the pt fails steroid therapy

What modality(ies) can be employed? Cryoablation, or laser photocoagulation

Which specific area should be treated?

If cryoablation is used, it should be applied...directly to the sclera adjacent to where snowbanking is present

If laser photocoagulation is used, it should be applied...to the retina adjacent to the snowbanking (but not to the snowbanking itself)

- 1) Steroid therapy
- 2) Peripheral retina ablation
- 3) Vitrectomy
- 4) Immunomodulatory therapy

- 1) The uveitis is profiled
- 2) The profiled case is meshed
- 3) A differential diagnosis list is generated
- 4) Studies are obtained to identify the etiology
- 5) Treatment appropriate for the etiology is initiated



Pars planitis

Intermediate uveitis

How is pars planitis/intermediate uveitis (PP/IU) managed?

When should peripheral retinal ablation be pursued?

If the pt fails steroid therapy

What modality(ies) can be employed?

Cryoablation, or laser photocoagulation

Which modality is preferred?

nere

If laser photocoagulation is used, it should be applied...to the retina adjacent to the snowbanking (but not to the snowbanking itself)

- 1) Steroid therapy
- 2) Peripheral retina ablation
- 3) Vitrectomy
- 4) Immunomodulatory therapy

- 1) The uveitis is profiled
- 2) The profiled case is meshed
- 3) A differential diagnosis list is generated
- 4) Studies are obtained to identify the etiology
- 5) Treatment appropriate for the etiology is initiated





Pars planitis

Intermediate uveitis



- 1) The uveitis is profiled
- 2) The profiled case is meshed
- 3) A differential diagnosis list is generated
- 4) Studies are obtained to identify the etiology
- 5) Treatment appropriate for the etiology is initiated



Pars planitis

Intermediate uveitis

How is pars planitis/intermediate uveitis (PP/IU) managed?

When should peripheral retinal ablation be pursued? If the pt fails steroid therapy

What modality(ies) can be employed?

Cryoablation, or laser photocoagulation

Which modality is preferred?

Probably laser photocoagulation. Cryoablation carries a risk of retinal detachment, which photocoagulation does not.

If laser photocoagulation is used, it should be applied...to the retina adjacent to the snowbanking (but not to the snowbanking itself)

- 1) Steroid therapy
- 2) Peripheral retina ablation
- 3) Vitrectomy
- 4) Immunomodulatory therapy

- 1) The uveitis is profiled
- 2) The profiled case is meshed
- 3) A differential diagnosis list is generated
- 4) Studies are obtained to identify the etiology
- 5) Treatment appropriate for the etiology is initiated





Intermediate uveitis

How is pars planitis/intermediate uveitis (PP/IU) managed? If an etiology is identified (ie, if it is IU), treatment specific to that etiology should be pursued

Assume testing is noncontributory. How should PP be managed?

If it is mild, and not causing significant morbidity, it can simply be monitored

Under what circumstances should treatment be initiated?

When should vitrectomy be pursued?

- 2) Peripheral retina ablation
- 3) Vitrectomy
- 4) Immunomodulatory therapy

- 1) The uveitis is profiled
- 2) The profiled case is meshed
- 3) A differential diagnosis list is generated
- 4) Studies are obtained to identify the etiology
- 5) Treatment appropriate for the etiology is initiated





Pars planitis

Intermediate uveitis

How is pars planitis/intermediate uveitis (PP/IU) managed? If an etiology is identified (ie, if it is IU), treatment specific to that etiology should be pursued

Assume testing is noncontributory. How should PP be managed? If it is mild, and not causing significant morbidity, it can simply be monitored

Under what circumstances should treatment be initiated?

When should vitrectomy be pursued?

If ablation fails to control the disease, and systemic immunomodulatory therapy is unacceptable

- Peripheral retina ablation
- 3) Vitrectomy
- 4) Immunomodulatory therapy

- 1) The uveitis is profiled
- 2) The profiled case is meshed
- 3) A differential diagnosis list is generated
- 4) Studies are obtained to identify the etiology
- 5) Treatment appropriate for the etiology is initiated



Pars planitis

Intermediate uveitis

If an etiology is identified (ie, if it is IU), treatment specific to that etiology should be pursued

Assume testing is noncontributory. How should PP be managed? If it is mild, and not causing significant morbidity, it can simply be monitored

When should vitrectomy be pursued?

If ablation fails to control the disease, and systemic immunomodulatory therapy is unacceptable

In addition to removal of the vitreous body, two other surgical maneuvers are desirable—what are they?

- pheral retina ablation
- 3) Vitrectomy
- emodulatory therapy

- 1) The uveitis is profiled
- 2) The profiled case is meshed
- 3) A differential diagnosis list is generated
- 4) Studies are obtained to identify the etiology
- 5) Treatment appropriate for the etiology is initiated



Pars planitis

Intermediate uveitis

How is pars planitis/intermediate uveitis (PP/IU) managed? If an etiology is identified (ie, if it is IU), treatment specific to that etiology should be pursued

Assume testing is noncontributory. How should PP be managed? If it is mild, and not causing significant morbidity, it can simply be monitored

Under what circumstances should treatment be initiated?

When should vitrectomy be pursued?

If ablation fails to control the disease, and systemic immunomodulatory therapy is unacceptable

In addition to removal of the vitreous body, two other surgical maneuvers are desirable—what are they?

Induction of a posterior vitreous detachment, and peripheral retinal photocoagulation

2) Peripheral retina ablation

3) Vitrectomy

4) Immune modulatory therapy

- 1) The uveitis is profiled
- 2) The profiled case is meshed
- 3) A differential diagnosis list is generated
- 4) Studies are obtained to identify the etiology
- 5) Treatment appropriate for the etiology is initiated



Pars planitis

Intermediate uveitis

How is pars planitis/intermediate uveitis (PP/IU) managed? If an etiology is identified (ie, if it is IU), treatment specific to that etiology should be pursued

Assume testing is noncontributory. How should PP be managed? If it is mild, and not causing significant morbidity, it can simply be monitored

When should immunomodulatory therapy be pursued?

3) Vitrectemy

4) Immunomodulatory therapy

- 1) The uveitis is profiled
- 2) The profiled case is meshed
- 3) A differential diagnosis list is generated
- 4) Studies are obtained to identify the etiology
- 5) Treatment appropriate for the etiology is initiated



Pars planitis

Intermediate uveitis

How is pars planitis/intermediate uveitis (PP/IU) managed? If an etiology is identified (ie, if it is IU), treatment specific to that etiology should be pursued

Assume testing is noncontributory. How should PP be managed? If it is mild, and not causing significant morbidity, it can simply be monitored

When should immunomodulatory therapy be pursued?

If other interventions failed, and/or if two words disease is present

3) Vitrectemy

4) Immunomodulatory therapy

- 1) The uveitis is profiled
- 2) The profiled case is meshed
- 3) A differential diagnosis list is generated
- 4) Studies are obtained to identify the etiology
- 5) Treatment appropriate for the etiology is initiated



Pars planitis

Intermediate uveitis

How is pars planitis/intermediate uveitis (PP/IU) managed? If an etiology is identified (ie, if it is IU), treatment specific to that etiology should be pursued

Assume testing is noncontributory. How should PP be managed? If it is mild, and not causing significant morbidity, it can simply be monitored

When should immunomodulatory therapy be pursued? If other interventions failed, and/or if severe bilateral disease is present

3) Vitrectomy

4) Immunomodulatory therapy

- 1) The uveitis is profiled
- 2) The profiled case is meshed
- 3) A differential diagnosis list is generated
- 4) Studies are obtained to identify the etiology
- 5) Treatment appropriate for the etiology is initiated



Pars planitis

Intermediate uveitis

How is pars planitis/intermediate uveitis (PP/IU) managed? If an etiology is identified (ie, if it is IU), treatment specific to that etiology should be pursued

Assume testing is noncontributory. How should PP be managed? If it is mild, and not causing significant morbidity, it can simply be monitored

When should immunomodulatory therapy be pursued?

If other interventions failed, and/or if severe bilateral disease is present

Which agents can/should be considered?

- --?
- --?
- --?
- --?
 - 3) Vitrectomy
 - 4) Immunomodulatory therapy

- 1) The uveitis is profiled
- 2) The profiled case is meshed
- 3) A differential diagnosis list is generated
- 4) Studies are obtained to identify the etiology
- 5) Treatment appropriate for the etiology is initiated



Pars planitis

Intermediate uveitis

How is pars planitis/intermediate uveitis (PP/IU) managed? If an etiology is identified (ie, if it is IU), treatment specific to that etiology should be pursued

Assume testing is noncontributory. How should PP be managed? If it is mild, and not causing significant morbidity, it can simply be monitored

When should immunomodulatory therapy be pursued?

If other interventions failed, and/or if severe bilateral disease is present

Which agents can/should be considered?

- --Methotrexate
- --Cyclosporine
- --Azathioprine
- -- Cyclophosphamide
 - 3) Vitrectemy
 - 4) Immunomodulatory therapy