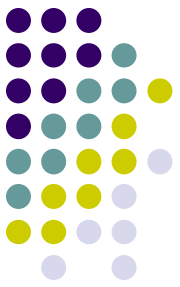


Q

Hyphema

What is the definition of hyphema?

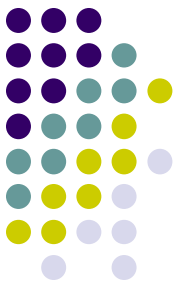


A

Hyphema

What is the definition of hyphema?

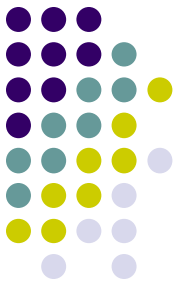
The presence of RBCs in the anterior chamber



Q

Hyphema

3



What is the definition of hyphema?

The presence of RBCs in the anterior chamber

With respect to its extent, there are two types of hyphema—what are they?

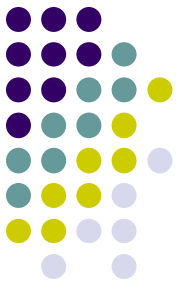
--

--

Q/A

Hyphema

4



What is the definition of hyphema?

The presence of RBCs in the anterior chamber

With respect to its extent, there are two types of hyphema—what are they?

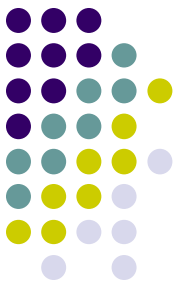
--When the extent of the hyphema is limited to a few RBCs circulating in the AC, it is called a

--When the hyphema is extensive enough to form a clot in the AC, it is called a hyphema

A

Hyphema

5



What is the definition of hyphema?

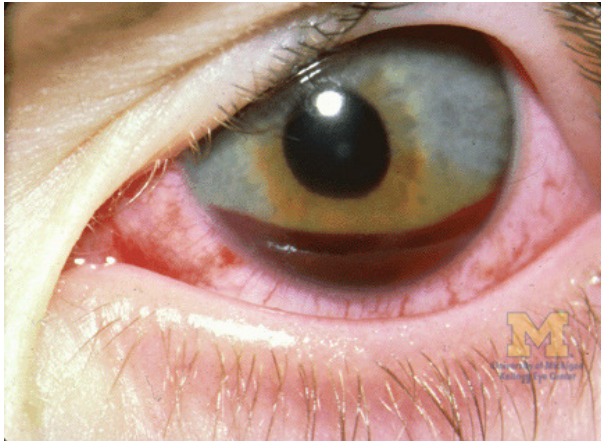
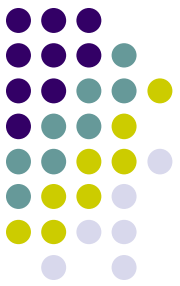
The presence of RBCs in the anterior chamber

With respect to its extent, there are two types of hyphema—what are they?

--When the extent of the hyphema is limited to a few RBCs circulating in the AC, it is called a microhyphema

--When the hyphema is extensive enough to form a clot in the AC, it is called a layered hyphema

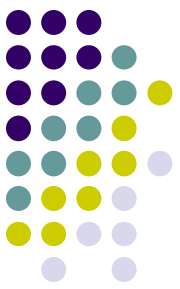
Hyphema



Layered hyphema

Q

Hyphema



What is the definition of hyphema?

The presence of RBCs in the anterior chamber

With respect to its extent, there are two types of hyphema—what are they?

--When the extent of the hyphema is limited to a few RBCs circulating in the AC, it is called a microhyphema

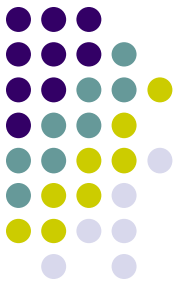
--When the hyphema is extensive enough to form a clot in the AC, it is called a layered hyphema

What is the most common cause of hyphema?

A

Hyphema

8



What is the definition of hyphema?

The presence of RBCs in the anterior chamber

With respect to its extent, there are two types of hyphema—what are they?

--When the extent of the hyphema is limited to a few RBCs circulating in the AC, it is called a microhyphema

--When the hyphema is extensive enough to form a clot in the AC, it is called a layered hyphema

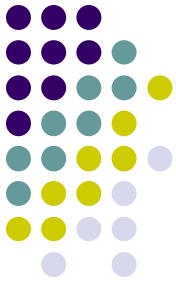
What is the most common cause of hyphema?

Trauma

Q

Hyphema

9



What is the definition of hyphema?

The presence of RBCs in the anterior chamber

With respect to its extent, there are two types of hyphema—what are they?

--When the extent of the hyphema is limited to a few RBCs circulating in the AC, it is called a microhyphema

--When the hyphema is extensive enough to form a clot in the AC, it is called a layered hyphema

What is the most common cause of hyphema?

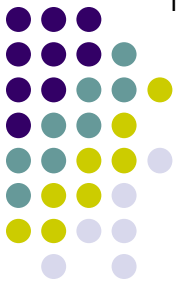
Trauma

What type of trauma is most commonly implicated?

A

Hyphema

10



What is the definition of hyphema?

The presence of RBCs in the anterior chamber

With respect to its extent, there are two types of hyphema—what are they?

--When the extent of the hyphema is limited to a few RBCs circulating in the AC, it is called a microhyphema

--When the hyphema is extensive enough to form a clot in the AC, it is called a layered hyphema

What is the most common cause of hyphema?

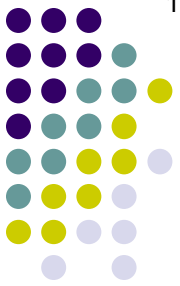
Trauma

What type of trauma is most commonly implicated?

Blunt

Q

Hyphema



What is the definition of hyphema?

The presence of RBCs in the anterior chamber

With respect to its extent, there are two types of hyphema—what are they?

--When the extent of the hyphema is limited to a few RBCs circulating in the AC, it is called a microhyphema

--When the hyphema is extensive enough to form a clot in the AC, it is called a layered hyphema

What is the most common cause of hyphema?

Trauma

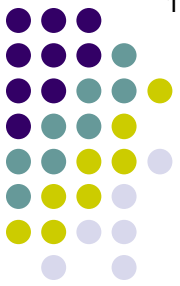
What is the mechanism by which blunt trauma causes hyphema?

What type

Blunt

A

Hyphema



What is the definition of hyphema?

The presence of RBCs in the anterior chamber

With respect to its extent, there are two types of hyphema—what are they?

--When the extent of the hyphema is limited to a few RBCs circulating in the AC, it is called a microhyphema

--When the hyphema is extensive enough to form a clot in the AC, it is called a layered hyphema

What is the most common cause of hyphema?

Trauma

What is the mechanism by which blunt trauma causes hyphema?

The globe is incompressible in the sense that its volume cannot be reduced. (It's like a balloon—squeeze it in one place, it has to expand in another.)

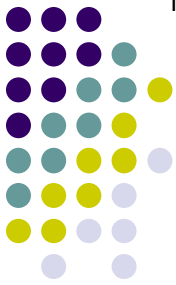
So when blunt force to the globe compresses its anterior-posterior dimension, the globe compensates by expanding in the equatorial plane—the eye gets momentarily shorter and fatter.

What type

Blunt

A

Hyphema



What is the definition of hyphema?

The presence of RBCs in the anterior chamber

With respect to its extent, there are two types of hyphema—what are they?

--When the extent of the hyphema is limited to a few RBCs circulating in the AC, it is called a microhyphema

--When the hyphema is extensive enough to form a clot in the AC, it is called a layered hyphema

What is the most common cause of hyphema?

Trauma

What is the mechanism by which blunt trauma causes hyphema?

The globe is incompressible in the sense that its volume cannot be reduced.

(It's like a balloon—squeeze it in one place, it has to expand in another.)

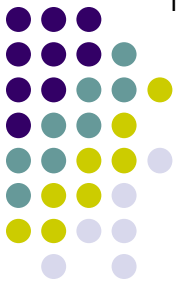
So when blunt force to the globe compresses its anterior-posterior dimension, the globe compensates by expanding in the equatorial plane—the eye gets momentarily shorter and fatter. In turn, equatorial expansion stretches the iris, including its major (and other) arterial circle. Tears of these vessels or their branches are the common source of blood in traumatic hyphema.

What type

Blunt



Hyphema



What is the definition of hyphema?

The presence of RBCs in the anterior chamber

With respect to its extent, there are two types of hyphema—what are they?

--When the extent of the hyphema is limited to a few RBCs circulating in the AC, it is called a microhyphema

--When the hyphema is extensive enough to form a clot in the AC, it is called a layered hyphema

What is the most common cause of hyphema?

Trauma

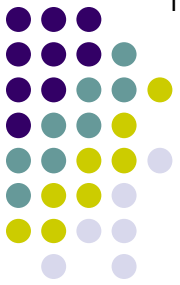
What type of trauma is most commonly implicated?

Blunt

Who is at risk for traumatic hyphema?

A

Hyphema



What is the definition of hyphema?

The presence of RBCs in the anterior chamber

With respect to its extent, there are two types of hyphema—what are they?

--When the extent of the hyphema is limited to a few RBCs circulating in the AC, it is called a microhyphema

--When the hyphema is extensive enough to form a clot in the AC, it is called a layered hyphema

What is the most common cause of hyphema?

Trauma

What type of trauma is most commonly implicated?

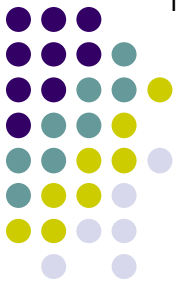
Blunt

Who is at risk for traumatic hyphema?

Young men

Q

Hyphema



What is the definition of hyphema?

The presence of RBCs in the anterior chamber

With respect to its extent, there are two types of hyphema—what are they?

--When the extent of the hyphema is limited to a few RBCs circulating in the AC, it is called a microhyphema

--When the hyphema is extensive enough to form a clot in the AC, it is called a layered hyphema

What is the most common cause of hyphema?

Trauma

What type of trauma is most commonly implicated?

Blunt

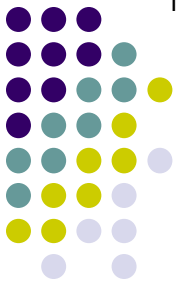
Who is at risk for traumatic hyphema?

Young men

Why are young men at risk?

A

Hyphema



What is the definition of hyphema?

The presence of RBCs in the anterior chamber

With respect to its extent, there are two types of hyphema—what are they?

--When the extent of the hyphema is limited to a few RBCs circulating in the AC, it is called a microhyphema

--When the hyphema is extensive enough to form a clot in the AC, it is called a layered hyphema

What is the most common cause of hyphema?

Trauma

What type of trauma is most commonly implicated?

Blunt

Who is at risk for traumatic hyphema?

Young men

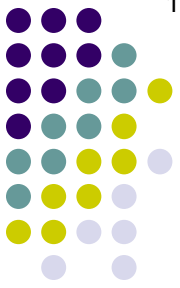
Why are young men at risk?

Because they're dumb, and do things that lead to getting hit in the eye

Q

Hyphema

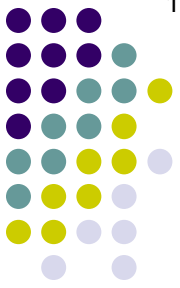
- What is the pre-eminent goal in managing hyphema?



A

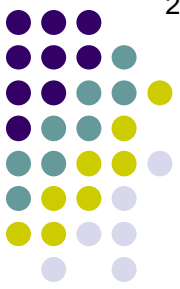
Hyphema

- What is the pre-eminent goal in managing hyphema? **Avoid a re-bleed**



Q

Hyphema

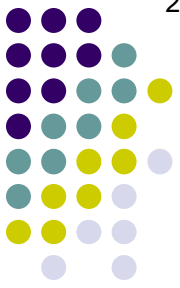


- What is the pre-eminent goal in managing hyphema? **Avoid a re-bleed**

Is re-bleeding a common event in hyphema?

Q/A

Hyphema



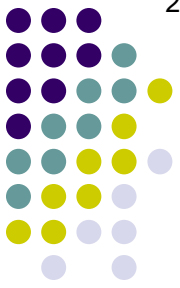
- What is the pre-eminent goal in managing hyphema? **Avoid a re-bleed**

Is re-bleeding a common event in hyphema?

Not really; only somewhere around % of cases re-bleed

A

Hyphema



- What is the pre-eminent goal in managing hyphema? **Avoid a re-bleed**

Is re-bleeding a common event in hyphema?

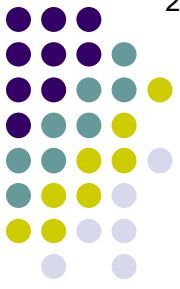
Not really; only somewhere around 5% of cases re-bleed

Re-bleed rate per:

--*Glaucoma* book: 5-10%

--*Cornea* book: <5%

Q

Hyphema

- What is the pre-eminent goal in managing hyphema? **Avoid a re-bleed**

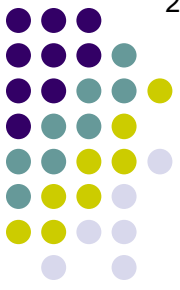
Is re-bleeding a common event in hyphema?

Not really; only somewhere around 5% of cases re-bleed

Relative to the initial bleed, when is a re-bleed likely to occur?

A

Hyphema



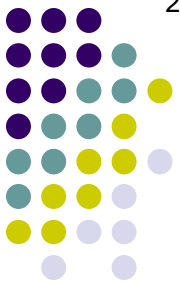
- What is the pre-eminent goal in managing hyphema? **Avoid a re-bleed**

Is re-bleeding a common event in hyphema?

Not really; only somewhere around 5% of cases re-bleed

Relative to the initial bleed, when is a re-bleed likely to occur?

3-7 days post-event



Q

Hyphema

- What is the pre-eminent goal in managing hyphema? **Avoid a re-bleed**

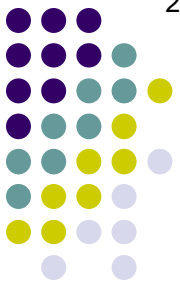
Is re-bleeding a common event in hyphema?

Not really; only somewhere around 5% of cases re-bleed

Relative to the initial bleed, when is a re-bleed likely to occur?

3-7 days post-event

Why then? What happens 3-7 days after the initial bleed?



A

Hyphema

- What is the pre-eminent goal in managing hyphema? **Avoid a re-bleed**

Is re-bleeding a common event in hyphema?

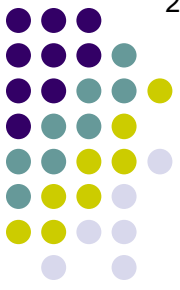
Not really; only somewhere around 5% of cases re-bleed

Relative to the initial bleed, when is a re-bleed likely to occur?

3-7 days post-event

Why then? What happens 3-7 days after the initial bleed?

This is when the original clot is going through the process of lysis/retraction



Q

Hyphema

- What is the pre-eminent goal in managing hyphema? **Avoid a re-bleed**

Is re-bleeding a common event in hyphema?

Not really; only somewhere around 5% of cases re-bleed

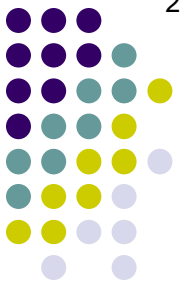
Relative to the initial bleed, when is a re-bleed likely to occur?

3-7 days post-event

What's the big deal about re-bleeding, ie, why is avoiding it so important?

A

Hyphema



- What is the pre-eminent goal in managing hyphema? **Avoid a re-bleed**

Is re-bleeding a common event in hyphema?

Not really; only somewhere around 5% of cases re-bleed

Relative to the initial bleed, when is a re-bleed likely to occur?

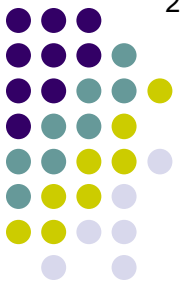
3-7 days post-event

What's the big deal about re-bleeding, ie, why is avoiding it so important?

Because the risk of long-term complications goes up significantly if re-bleeding occurs

Q

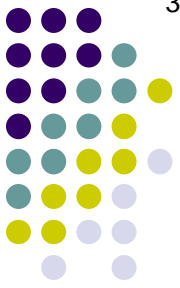
Hyphema



- What is the pre-eminent goal in managing hyphema? **Avoid a re-bleed**
- What goal is a close second?

A

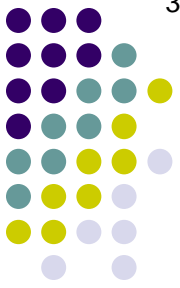
Hyphema



- What is the pre-eminent goal in managing hyphema? **Avoid a re-bleed**
- What goal is a close second? **Control IOP**

Q

Hyphema

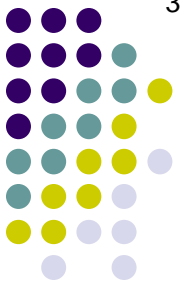


- What is the pre-eminent goal in managing hyphema? Avoid a re-bleed
- What goal is a close second? **Control IOP**

What is the mechanism by which hyphema leads to elevated IOP?

A

Hyphema

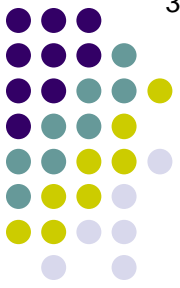


- What is the pre-eminent goal in managing hyphema? Avoid a re-bleed
- What goal is a close second? **Control IOP**

What is the mechanism by which hyphema leads to elevated IOP?

The TM becomes clogged with RBCs, along with the usual inflammatory cells and material.
(Not to mention, the inciting trauma may have damaged the angle.)

Q

Hyphema

- What is the pre-eminent goal in managing hyphema? Avoid a re-bleed
- What goal is a close second? **Control IOP**

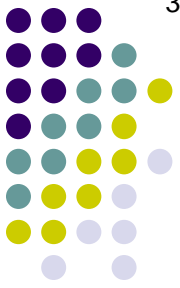
What is the mechanism by which hyphema leads to elevated IOP?

The TM becomes clogged with RBCs, along with the usual inflammatory cells and material.
(Not to mention, the inciting trauma may have damaged the angle.)

What event modestly increases the risk of significant IOP elevation?

A

Hyphema



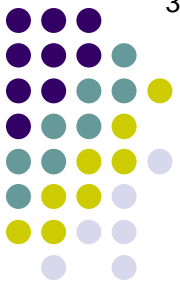
- What is the pre-eminent goal in managing hyphema? Avoid a re-bleed
- What goal is a close second? **Control IOP**

What is the mechanism by which hyphema leads to elevated IOP?

The TM becomes clogged with RBCs, along with the usual inflammatory cells and material.
(Not to mention, the inciting trauma may have damaged the angle.)

What event modestly increases the risk of significant IOP elevation?

A large hyphema (ie, there is a modest correlation between hyphema size and the incidence of IOP elevation)



Q

Hyphema

- What is the pre-eminent goal in managing hyphema? Avoid a re-bleed
- What goal is a close second? **Control IOP**

What is the mechanism by which hyphema leads to elevated IOP?

The TM becomes clogged with RBCs, along with the usual inflammatory cells and material.
(Not to mention, the inciting trauma may have damaged the angle.)

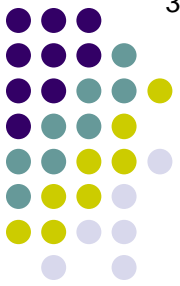
What event modestly increases the risk of significant IOP elevation?

A large hyphema (ie, there is a modest correlation between hyphema size and the incidence of IOP elevation)

*What event **greatly** increases the risk of significant IOP elevation?*

A

Hyphema



- What is the pre-eminent goal in managing hyphema? Avoid a re-bleed
- What goal is a close second? **Control IOP**

What is the mechanism by which hyphema leads to elevated IOP?

The TM becomes clogged with RBCs, along with the usual inflammatory cells and material.
(Not to mention, the inciting trauma may have damaged the angle.)

What event modestly increases the risk of significant IOP elevation?

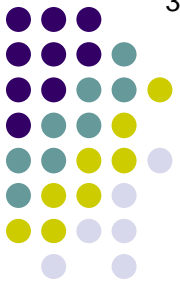
A large hyphema (ie, there is a modest correlation between hyphema size and the incidence of IOP elevation)

*What event **greatly** increases the risk of significant IOP elevation?*

A re-bleed

Q

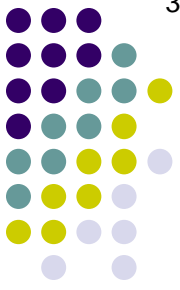
Hyphema



- What is the pre-eminent goal in managing hyphema? **Avoid a re-bleed**
- What goal is a close second? **Control IOP**
- Third?

A

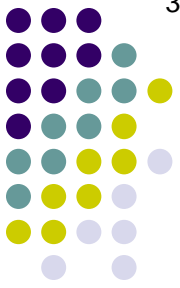
Hyphema



- What is the pre-eminent goal in managing hyphema? **Avoid a re-bleed**
- What goal is a close second? **Control IOP**
- Third? **Control inflammation**

Q

Hyphema

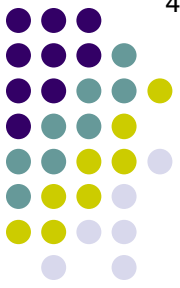


- What is the pre-eminent goal in managing hyphema? Avoid a re-bleed
- What goal is a close second? Control IOP
- Third? **Control inflammation**

Why is inflammation control important?

A

Hyphema

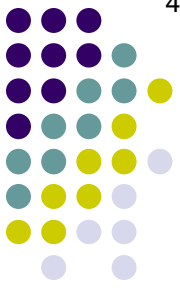


- What is the pre-eminent goal in managing hyphema? Avoid a re-bleed
- What goal is a close second? Control IOP
- Third? **Control inflammation**

Why is inflammation control important?

To reduce the risk of synechiae formation (and to improve pt comfort of course).

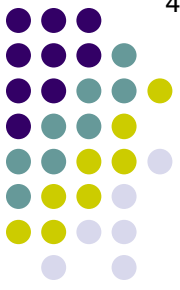
Q

Hyphema

- What is the pre-eminent goal in managing hyphema? **Avoid a re-bleed**
- What goal is a close second? **Control IOP**
- Third? **Control inflammation**
- There is another goal that, in young children, is arguably as important as avoiding re-bleed. What is it?

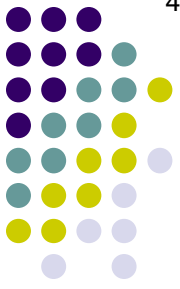
A

Hyphema



- What is the pre-eminent goal in managing hyphema? **Avoid a re-bleed**
- What goal is a close second? **Control IOP**
- Third? **Control inflammation**
- There is another goal that, in young children, is arguably as important as avoiding re-bleed. What is it? **Prevent corneal bloodstaining**

Q

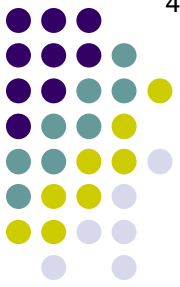
Hyphema

- What is the pre-eminent goal in managing hyphema? Avoid a re-bleed
- What goal is a close second? Control IOP
- Third? Control inflammation
- There is another goal that, in young children, is arguably as important as avoiding re-bleed. What is it? **Prevent corneal bloodstaining**

Briefly, how does a hyphema lead to corneal bloodstaining?

Q/A

Hyphema



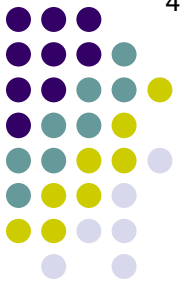
- What is the pre-eminent goal in managing hyphema? Avoid a re-bleed
- What goal is a close second? Control IOP
- Third? Control inflammation
- There is another goal that, in young children, is arguably as important as avoiding re-bleed. What is it? **Prevent corneal bloodstaining**

Briefly, how does a hyphema lead to corneal bloodstaining?

RBCs in the AC release abb, which enters the corneal stroma and gets absorbed by keratocytes

A

Hyphema

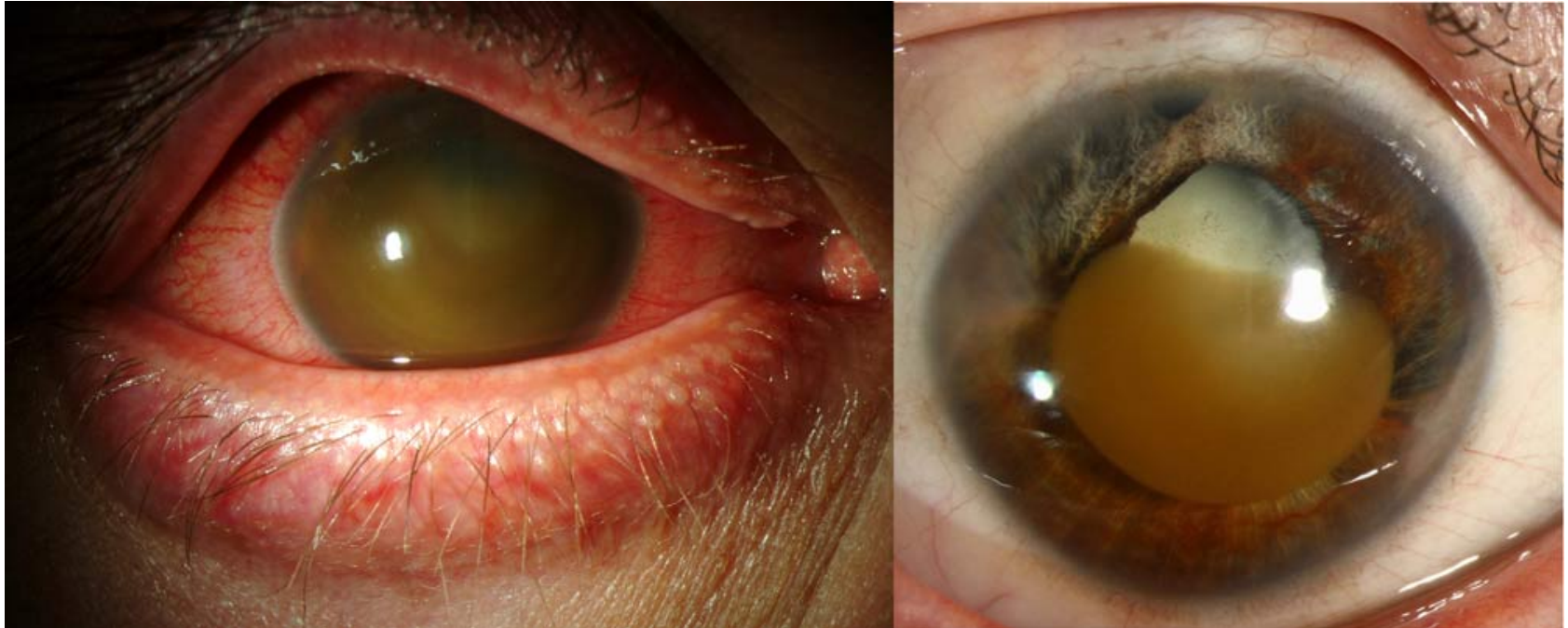
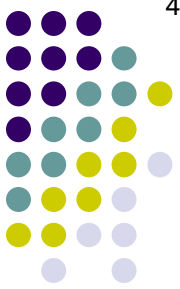


- What is the pre-eminent goal in managing hyphema? Avoid a re-bleed
- What goal is a close second? Control IOP
- Third? Control inflammation
- There is another goal that, in young children, is arguably as important as avoiding re-bleed. What is it? **Prevent corneal bloodstaining**

Briefly, how does a hyphema lead to corneal bloodstaining?

RBCs in the AC release Hgb, which enters the corneal stroma and gets absorbed by keratocytes

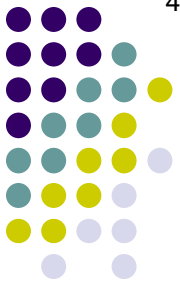
Hyphema



Corneal bloodstaining

A

Hyphema



- What is the pre-eminent goal in managing hyphema? Avoid a re-bleed
- What goal is a close second? Control IOP
- Third? Control inflammation
- There is another goal that, in young children, is arguably as important as avoiding re-bleed. What is it? **Prevent corneal bloodstaining**

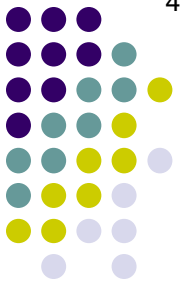
Briefly, how does a hyphema lead to corneal bloodstaining?

RBCs in the AC release Hgb, which enters the corneal stroma and gets absorbed by keratocytes

What complication—common in hyphema—increases the risk of bloodstaining?

A

Hyphema



- What is the pre-eminent goal in managing hyphema? Avoid a re-bleed
- What goal is a close second? Control IOP
- Third? Control inflammation
- There is another goal that, in young children, is arguably as important as avoiding re-bleed. What is it? **Prevent corneal bloodstaining**

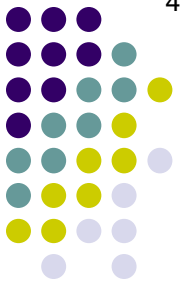
Briefly, how does a hyphema lead to corneal bloodstaining?

RBCs in the AC release Hgb, which enters the corneal stroma and gets absorbed by keratocytes

What complication—common in hyphema—increases the risk of bloodstaining?

Increased IOP

Q

Hyphema

- What is the pre-eminent goal in managing hyphema? Avoid a re-bleed
- What goal is a close second? Control IOP
- Third? Control inflammation
- There is another goal that, in young children, is arguably as important as avoiding re-bleed. What is it? **Prevent corneal bloodstaining**

Briefly, how does a hyphema lead to corneal bloodstaining?

RBCs in the AC release Hgb, which enters the corneal stroma and gets absorbed by keratocytes

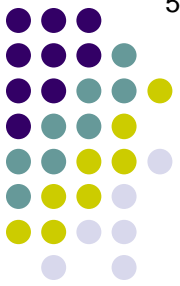
What complication—common in hyphema—increases the risk of bloodstaining?

Increased IOP

Why is corneal bloodstaining such a concern in young children?

A

Hyphema



- What is the pre-eminent goal in managing hyphema? Avoid a re-bleed
- What goal is a close second? Control IOP
- Third? Control inflammation
- There is another goal that, in young children, is arguably as important as avoiding re-bleed. What is it? **Prevent corneal bloodstaining**

Briefly, how does a hyphema lead to corneal bloodstaining?

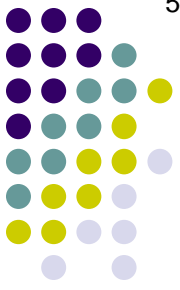
RBCs in the AC release Hgb, which enters the corneal stroma and gets absorbed by keratocytes

What complication—common in hyphema—increases the risk of bloodstaining?

Increased IOP

Why is corneal bloodstaining such a concern in young children?

Because it is potentially amblyogenic



Q

Hyphema

- What is the pre-eminent goal in managing hyphema? Avoid a re-bleed
- What goal is a close second? Control IOP
- Third? Control inflammation
- There is another goal that, in young children, is arguably as important as avoiding re-bleed. What is it? **Prevent corneal bloodstaining**

Briefly, how does a hyphema lead to corneal bloodstaining?

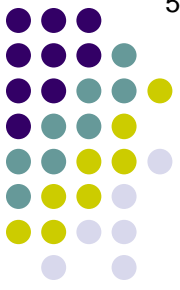
RBCs in the AC release hemoglobin, which is then absorbed by keratocytes. *Is the prognosis for corneal bloodstaining good, or bad?*

What complication—common in hyphema—increases the risk of bloodstaining?
Increased IOP

Why is corneal bloodstaining such a concern in young children?
Because it is potentially amblyogenic

A

Hyphema



- What is the pre-eminent goal in managing hyphema? Avoid a re-bleed
- What goal is a close second? Control IOP
- Third? Control inflammation
- There is another goal that, in young children, is arguably as important as avoiding re-bleed. What is it? **Prevent corneal bloodstaining**

Briefly, how does a hyphema lead to corneal bloodstaining?

RBCs in the AC release hemoglobin, which is then absorbed by keratocytes.

Is the prognosis for corneal bloodstaining good, or bad?

Both. It spontaneously and completely clears (yay!) over a course of months to years (boo!)

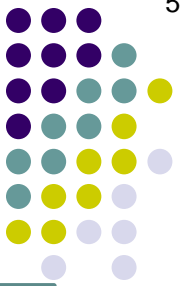
What complication—common in hyphema—increases the risk of bloodstaining?

Increased IOP

Why is corneal bloodstaining such a concern in young children?

Because it is potentially amblyogenic

Hyphema



Avoid a re-bleed

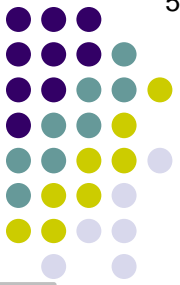
Control IOP

Control inflammation

Prevent corneal bloodstaining

tl;dr The treatment goals in managing hyphema

No question—proceed when ready

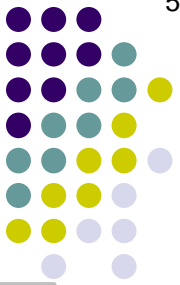


Avoid a re-bleed

Speaking of managing hyphema... What steps should be taken to minimize the risk of re-bleeding?

- three words around the clock
- two/three different words around the clock
- Strict
- Avoidance of
- See the pt unit of time until the hyphema resolves

tl;dr The treatment goals in managing hyphema



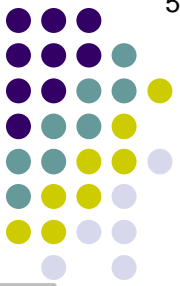
Avoid a re-bleed

Speaking of managing hyphema...What steps should be taken to minimize the risk of re-bleeding?

- Shield the eye around the clock
- Elevate the head around the clock
- Strict bedrest
- Avoidance of anticoagulants
- See the pt daily until the hyphema resolves

Prevent corneal bloodstaining

tl;dr The treatment goals in managing hyphema



Avoid a re-bleed

Speaking of managing hyphema... What steps should be taken to minimize the risk of re-bleeding?

-- Shield the eye around the clock

-- Elevate the head and limit activity

-- **Strict bedrest**

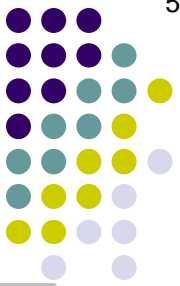
Does this mean the pt should be hospitalized?

-- Avoidance of anticoagulants

-- See the pt daily until the hyphema resolves

Prevent corneal bloodstaining

tl;dr The treatment goals in managing hyphema



Avoid a re-bleed

Speaking of managing hyphema... What steps should be taken to minimize the risk of re-bleeding?

-- Shield the eye around the clock

-- Elevate the head around the clock

-- **Strict bedrest**

Does this mean the pt should be hospitalized?

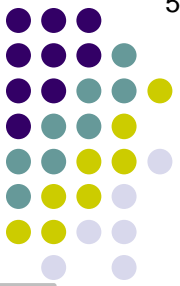
If necessary, yes (as is often the case with)

-- Avoidance of anticoagulants

-- See the pt daily until the hyphema resolves

Prevent corneal bloodstaining

tl;dr The treatment goals in managing hyphema



Avoid a re-bleed

Speaking of managing hyphema... What steps should be taken to minimize the risk of re-bleeding?

-- Shield the eye around the clock

-- Elevate the head around the clock

-- **Strict bedrest**

Does this mean the pt should be hospitalized?

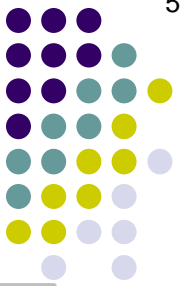
If necessary, yes (as is often the case with children)

-- Avoidance of anticoagulants

-- See the pt daily until the hyphema resolves

Prevent corneal bloodstaining

tl;dr The treatment goals in managing hyphema



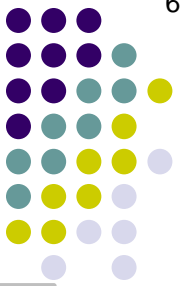
Avoid a re-bleed

Speaking of managing hyphema... What steps should be taken to minimize the risk of re-bleeding?

- Shield the eye around the clock
- Elevate the head around the clock
- Strict bedrest
- **Avoidance of anticoagulants**
- See the pt daily until the hyphema resolves

How might a pt inadvertently anti-coagulate him/herself?

tl;dr The treatment goals in managing hyphema



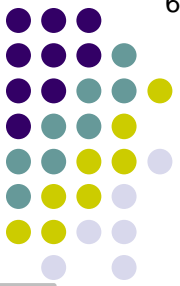
Avoid a re-bleed

Speaking of managing hyphema... What steps should be taken to minimize the risk of re-bleeding?

- Shield the eye around the clock
- Elevate the head around the clock
- Strict bedrest
- **Avoidance of anticoagulants**
- See the pt daily until the hyphema resolves

How might a pt inadvertently anti-coagulate him/herself?
By taking aspirin for pain

tl;dr The treatment goals in managing hyphema



Avoid a re-bleed

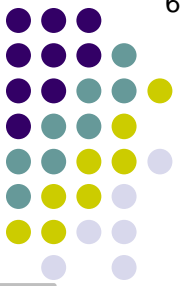
Speaking of managing hyphema... What steps should be taken to minimize the risk of re-bleeding?

- Shield the eye around the clock
- Elevate the head around the clock
- Strict bedrest
- **Avoidance of anticoagulants**
- See the pt daily until the hyphema resolves

How might a pt inadvertently anti-coagulate him/herself?
By taking aspirin for pain

What should you tell the pt in this regard?

tl;dr The treatment goals in managing hyphema



Avoid a re-bleed

Speaking of managing hyphema... What steps should be taken to minimize the risk of re-bleeding?

- Shield the eye around the clock
- Elevate the head around the clock
- Strict bedrest
- **Avoidance of anticoagulants**
- See the pt daily until the hyphema resolves

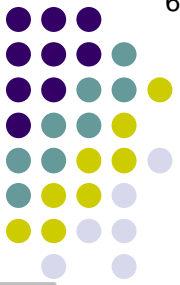
How might a pt inadvertently anti-coagulate him/herself?
By taking aspirin for pain

What should you tell the pt in this regard?
To avoid aspirin (and other NSAIDs, just to be safe)

tl;dr The treatment goals in managing hyphema

Q

Hyphema

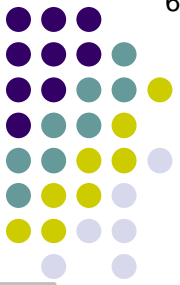


Avoid a re-bleed

Speaking of managing hyphema... What steps should be taken to minimize the risk of re-bleeding?

-- Shield the eye around the clock

What is aminocaproic acid, and how does it relate to avoiding re-bleed in hyphema management?



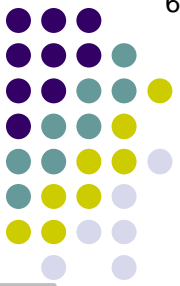
Avoid a re-bleed

Speaking of managing hyphema... What steps should be taken to minimize the risk of re-bleeding?

-- Shield the eye around the clock

What is aminocaproic acid, and how does it relate to avoiding re-bleed in hyphema management?

It is a systemic med that acts as a 'clot stabilizer' by enhancing hemostasis during the process of clot fibrinolysis



Avoid a re-bleed

Speaking of managing hyphema... What steps should be taken to minimize the risk of re-bleeding?

-- Shield the eye around the clock

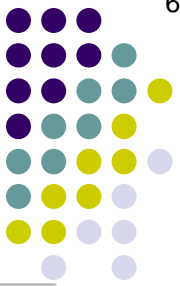
What is aminocaproic acid, and how does it relate to avoiding re-bleed in hyphema management?

It is a systemic med that acts as a 'clot stabilizer' by enhancing hemostasis during the process of clot fibrinolysis

What is the rationale behind its use in managing hyphema?

Q

Hyphema



Avoid a re-bleed

Speaking of managing hyphema... What steps should be taken to minimize the risk of re-bleeding?

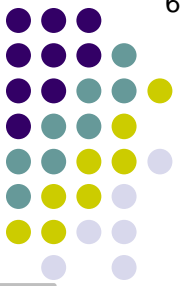
-- Shield the eye around the clock

What is aminocaproic acid, and how does it relate to avoiding re-bleed in hyphema management?

It is a systemic med that acts as a 'clot stabilizer' by enhancing hemostasis during the process of clot fibrinolysis

What is the rationale behind its use in managing hyphema?

By enhancing hemostasis, it may reduce the risk of a re-bleed



Avoid a re-bleed

Speaking of managing hyphema... What steps should be taken to minimize the risk of re-bleeding?

-- Shield the eye around the clock

What is aminocaproic acid, and how does it relate to avoiding re-bleed in hyphema management?

It is a systemic med that acts as a 'clot stabilizer' by enhancing hemostasis during the process of clot fibrinolysis

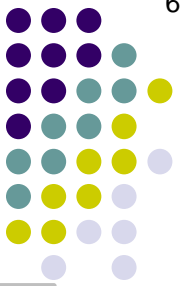
What is the rationale behind its use in managing hyphema?

By enhancing hemostasis, it may reduce the risk of a re-bleed

Aminocaproic acid is not often used in hyphema management—why?

Q

Hyphema



Avoid a re-bleed

Speaking of managing hyphema... What steps should be taken to minimize the risk of re-bleeding?

-- Shield the eye around the clock

What is aminocaproic acid, and how does it relate to avoiding re-bleed in hyphema management?

It is a systemic med that acts as a 'clot stabilizer' by enhancing hemostasis during the process of clot fibrinolysis

What is the rationale behind its use in managing hyphema?

By enhancing hemostasis, it may reduce the risk of a re-bleed

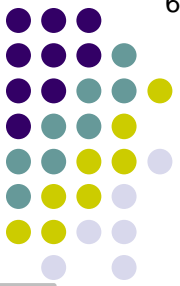
Aminocaproic acid is not often used in hyphema management—why?

Three reasons:

--

--

--



Avoid a re-bleed

Speaking of managing hyphema... What steps should be taken to minimize the risk of re-bleeding?

-- Shield the eye around the clock

What is aminocaproic acid, and how does it relate to avoiding re-bleed in hyphema management?

It is a systemic med that acts as a 'clot stabilizer' by enhancing hemostasis during the process of clot fibrinolysis

What is the rationale behind its use in managing hyphema?

By enhancing hemostasis, it may reduce the risk of a re-bleed

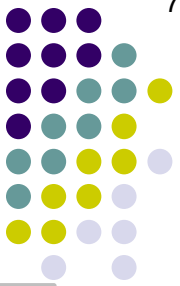
Aminocaproic acid is not often used in hyphema management—why?

Three reasons:

--The evidence regarding its ability to reduce re-bleed risk is equivocal

--It has myriad unwelcome side effects that limit its acceptance

--There is some evidence that the risk of re-bleed goes up when it is stopped



Avoid a re-bleed

Speaking of managing hyphema... What steps should be taken to minimize the risk of re-bleeding?

-- Shield the eye around the clock

What is aminocaproic acid, and how does it relate to avoiding re-bleed in hyphema management?

It is a systemic med that acts as a 'clot stabilizer' by enhancing hemostasis during the process of clot fibrinolysis

What is the rationale behind its use in managing hyphema?

By enhancing hemostasis, it may reduce the risk of a re-bleed

Aminocaproic acid is not often used in hyphema management—why?

Three reasons:

--The evidence regarding its ability to reduce

--It has **myriad unwelcome side effects** that

--There is some evidence that the risk of re-b

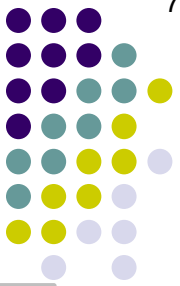
What are some of the myriad side effects?

--

--

--

--(There are many others)



Avoid a re-bleed

Speaking of managing hyphema... What steps should be taken to minimize the risk of re-bleeding?

-- Shield the eye around the clock

What is aminocaproic acid, and how does it relate to avoiding re-bleed in hyphema management?

It is a systemic med that acts as a 'clot stabilizer' by enhancing hemostasis during the process of clot fibrinolysis

What is the rationale behind its use in managing hyphema?

By enhancing hemostasis, it may reduce the risk of a re-bleed

Aminocaproic acid is not often used in hyphema management—why?

Three reasons:

--The evidence regarding its ability to reduce

--It has **myriad unwelcome side effects**

--There is some evidence that the risk of re-b

What are some of the myriad side effects?

--GI upset

--Hypotension (to the point of syncope)

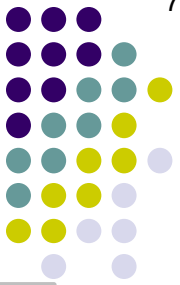
--Confusion

--(There are many others)

Q

Hyphema

72



Avoid a re-bleed

Control IOP

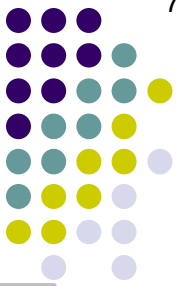
Control inflammation

Which two classes of meds are first-line in managing IOP in hyphema?

--

--

What are the treatment goals in managing hyphema?



Avoid a re-bleed

Control IOP

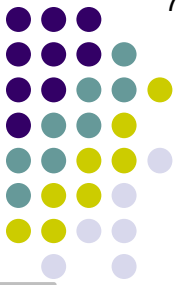
Control inflammation

Which two classes of meds are first-line in managing IOP in hyphema?

--Aqueous suppressants

--Hyperosmotic agents

What are the treatment goals in managing hyphema?



Avoid a re-bleed

Control IOP

Control inflammation

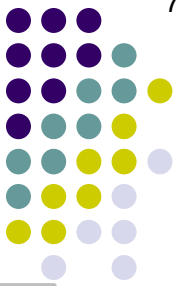
Which two classes of meds are first-line in managing IOP in hyphema?

Which three drop classes are aqueous suppressants?

--Aqueous suppressants?

--Hyperosmotic agents

What are the treatment goals in managing hyphema



Avoid a re-bleed

Control IOP

Control inflammation

Which two classes of meds are first-line in managing IOP in hyphema?

Which three drop classes are aqueous suppressants?

--Aqueous suppressants?

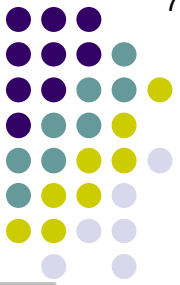
---- β blockers

---- α_2 agonists

----Carbonic anhydrase inhibitors (CAIs)

--Hyperosmotic agents

What are the treatment goals in managing hyphema?



Avoid a re-bleed

Control IOP

Control inflammation

Which two classes of meds are first-line in managing IOP in hyphema?

Which three drop classes are aqueous suppressants?

Which two meds are α_2 agonists?

--Aqueous suppressants

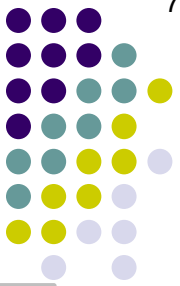
---- β blockers

---- α_2 **agonists?**

----Carbonic anhydrase inhibitors (CAIs)

--Hyperosmotic agents

What are the treatment goals in managing hyphema



Avoid a re-bleed

Control IOP

Control inflammation

Which two classes of meds are first-line in managing IOP in hyphema?

Which three drop classes are aqueous suppressants?

Which two meds are α_2 agonists?

Which two agents are hyperosmotics?

--Aqueous suppressants

---- β blockers

---- α_2 **agonists?**

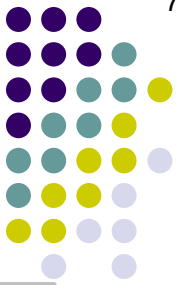
-----Apraclonidine (iopidine)

-----Brimonidine

----Carbonic anhydrase inhibitors (CAIs)

--Hyperosmotic agents

4, of the treatment goals in managing hyphema



Avoid a re-bleed

Control IOP

Control inflammation

Which two classes of meds are first-line in managing IOP in hyphema?

Which three drop classes are aqueous suppressants?

Which two meds are α_2 agonists?

Which two agents are hyperosmotics?

--Aqueous suppressants

---- β blockers

---- α_2 agonists

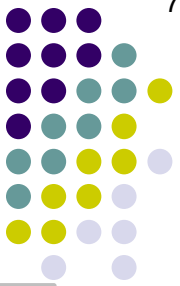
-----Apraclonidine (iopidine)

-----Brimonidine

----Carbonic anhydrase inhibitors (CAIs)

--**Hyperosmotic agents?**

What are the treatment goals in managing hyphema?



Avoid a re-bleed

Control IOP

Control inflammation

Which two classes of meds are first-line in managing IOP in hyphema?

Which three drop classes are aqueous suppressants?

Which two meds are α_2 agonists?

Which two agents are hyperosmotics?

--Aqueous suppressants

---- β blockers

---- α_2 agonists

-----Apraclonidine (iopidine)

-----Brimonidine

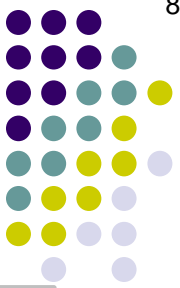
----Carbonic anhydrase inhibitors (CAIs)

--**Hyperosmotic agents?**

----Mannitol

----Glycerol

What are the treatment goals in managing hyphema?



Avoid a re-bleed

Control IOP

Control inflammation

Which two classes of meds are first-line in managing IOP in hyphema?

Which three drop classes are aqueous suppressants?

Which two meds are α_2 agonists?

Which two agents are hyperosmotics?

--Aqueous suppressants

If IOP can't be controlled medically, what surgical procedure should be considered?

-----Apraclonidine (iopidine)

-----Brimonidine

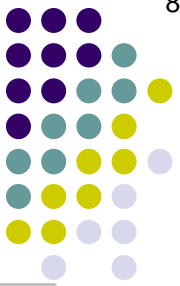
----Carbonic anhydrase inhibitors (CAIs)

--Hyperosmotic agents

----Mannitol

----Glycerol

What are the treatment goals in managing hyphema?



Avoid a re-bleed

Control IOP

Control inflammation

Which two classes of meds are first-line in managing IOP in hyphema?

Which three drop classes are aqueous suppressants?

Which two meds are α_2 agonists?

Which two agents are hyperosmotics?

--Aqueous suppressants

If IOP can't be controlled medically, what surgical procedure should be considered?
AC washout (we'll have more to say about this procedure later in the slide-set)

-----Apraclonidine (iopidine)

-----Brimonidine

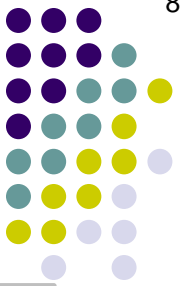
----Carbonic anhydrase inhibitors (CAIs)

--Hyperosmotic agents

----Mannitol

----Glycerol

What are the treatment goals in managing hyphema?



Avoid a re-bleed

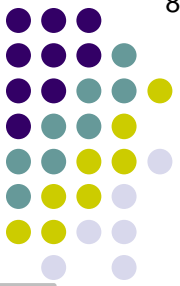
Control IOP

Control inflammation

How is intraocular inflammation controlled?

staining

tl;dr The treatment goals in managing hyphema



Avoid a re-bleed

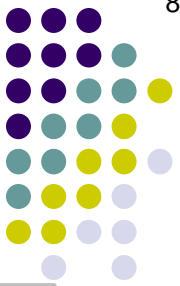
Control IOP

Control inflammation

How is intraocular inflammation controlled?
The usual way—with topical steroids and cycloplegics

staining

tl;dr The treatment goals in managing hyphema



Avoid a re-bleed

Control IOP

Control inflammation

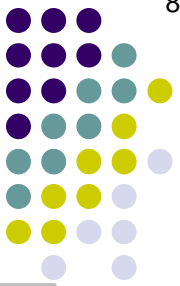
How is intraocular inflammation controlled?

The usual way—with topical steroids and cycloplegics

Note: Some clinicians forego steroids and cycloplegics in young children, contending that the potential benefit is outweighed by the risk incurred in rasslin' a struggling child to get the drops into her eye.

staining

tl;dr The treatment goals in managing hyphema



Avoid a re-bleed

Control IOP

Control inflammation

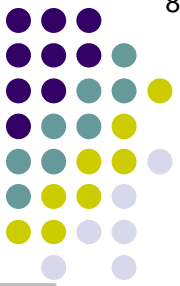
Prevent corneal bloodstaining

What steps can be taken to reduce the risk of corneal bloodstaining?

--

--

tl;dr The treatment goals in managing hyphema



Avoid a re-bleed

Control IOP

Control inflammation

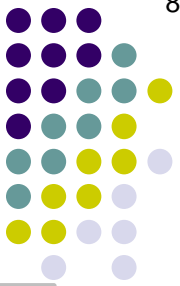
Prevent corneal bloodstaining

What steps can be taken to reduce the risk of corneal bloodstaining?

--Don't let IOP get out of hand

--Be vigilant clinically in assessing for evidence of bloodstaining, and have a low threshold for intervening surgically if necessary to prevent it from becoming visually significant

tl;dr The treatment goals in managing hyphema



Avoid a re-bleed

Control IOP

Control inflammation

Prevent corneal bloodstaining

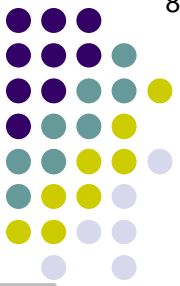
What steps can be taken to reduce the risk of corneal bloodstaining?

--Don't let IOP get out of hand

--Be vigilant clinically in assessing for evidence of bloodstaining, and have a low threshold for **intervening surgically** if necessary to prevent it from becoming visually significant

tl;dr The treatment

What surgical procedure are we talking about here?



Avoid a re-bleed

Control IOP

Control inflammation

Prevent corneal bloodstaining

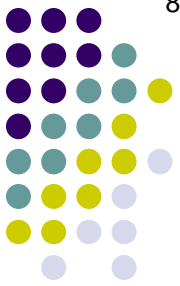
What steps can be taken to reduce the risk of corneal bloodstaining?

--Don't let IOP get out of hand

--Be vigilant clinically in assessing for evidence of bloodstaining, and have a low threshold for **intervening surgically** if necessary to prevent it from becoming visually significant

tl;dr The treatment

What surgical procedure are we talking about here?
Again, AC washout. **Speaking of...**



Q

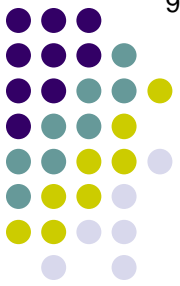
Hyphema

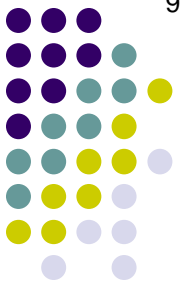
- *Hyphema: AC washout*
 - Goal is to remove two words from AC,
not the two different words

A

Hyphema

- *Hyphema: AC washout*
 - Goal is to remove **circulating RBCs** from AC, not the **entire clot**

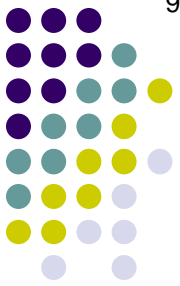




Q

Hyphema

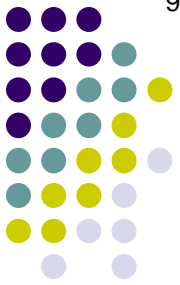
- *Hyphema: AC washout*
 - Goal is to remove **circulating RBCs** from AC, not the **entire clot**
 - Criteria triggering AC washout:
 - Any sign of **two words**



A

Hyphema

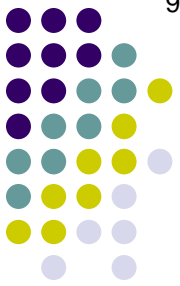
- *Hyphema: AC washout*
 - Goal is to remove **circulating RBCs** from AC, not the **entire clot**
 - Criteria triggering AC washout:
 - Any sign of **corneal bloodstaining**



Q

Hyphema

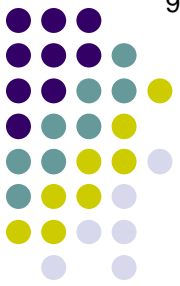
- *Hyphema: AC washout*
 - Goal is to remove **circulating RBCs** from AC, not the **entire clot**
 - Criteria triggering AC washout:
 - Any sign of **corneal bloodstaining**
 - Total hyphema x **amount of time**



A

Hyphema

- *Hyphema: AC washout*
 - Goal is to remove **circulating RBCs** from AC, not the **entire clot**
 - Criteria triggering AC washout:
 - Any sign of **corneal bloodstaining**
 - Total hyphema x **5 days**

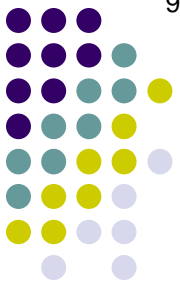


Q

Hyphema

- *Hyphema: AC washout*
 - Goal is to remove circulating RBCs from AC, not the entire clot
 - Criteria triggering AC washout:
 - Any sign of corneal bloodstaining
 - **Total hyphema x 5 days**

What complication is likely to occur if a total hyphema is present for more than 5 days?



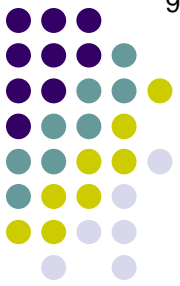
A

Hyphema

- *Hyphema: AC washout*
 - Goal is to remove circulating RBCs from AC, not the entire clot
 - Criteria triggering AC washout:
 - Any sign of corneal bloodstaining
 - **Total hyphema x 5 days**

What complication is likely to occur if a total hyphema is present for more than 5 days?

Peripheral anterior synechiae

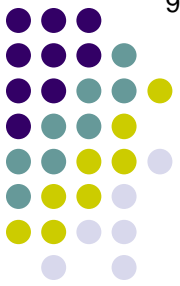


Q

Hyphema

- *Hyphema: AC washout*
 - Goal is to remove circulating RBCs from AC, not the entire clot
 - Criteria triggering AC washout:
 - Any sign of corneal bloodstaining
 - **Total hyphema x 5 days**

If a total hyphema consists of dark red-black blood, by name is it known?



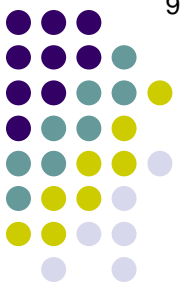
A

Hyphema

- *Hyphema: AC washout*
 - Goal is to remove circulating RBCs from AC, not the entire clot
 - Criteria triggering AC washout:
 - Any sign of corneal bloodstaining
 - **Total hyphema x 5 days**

*If a total hyphema consists of dark red-black blood,
by name is it known?
An 8-ball hyphema*

Hyphema



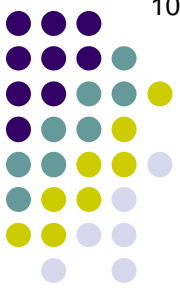
8-ball hyphema



Q

Hyphema

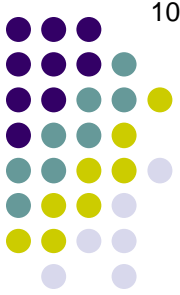
- *Hyphema: AC washout*
 - Goal is to remove **circulating RBCs** from AC, not the **entire clot**
 - Criteria triggering AC washout:
 - Any sign of **corneal bloodstaining**
 - Total hyphema x **5 days**
 - Hyphema >50% x **amount of time**



A

Hyphema

- *Hyphema: AC washout*
 - Goal is to remove **circulating RBCs** from AC, not the **entire clot**
 - Criteria triggering AC washout:
 - Any sign of **corneal bloodstaining**
 - Total hyphema x **5 days**
 - Hyphema >50% x **8 days**

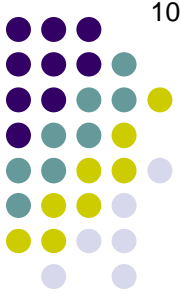


Q

Hyphema

- *Hyphema: AC washout*
 - Goal is to remove circulating RBCs from AC, not the entire clot
 - Criteria triggering AC washout:
 - Any sign of corneal bloodstaining
 - Total hyphema x 5 days
 - **Hyphema >50% x 8 days**

What complication is likely to occur if the IOP is >25 for longer than 5 days?



A

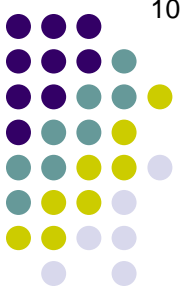
Hyphema

- *Hyphema: AC washout*
 - Goal is to remove circulating RBCs from AC, not the entire clot
 - Criteria triggering AC washout:
 - Any sign of corneal bloodstaining
 - Total hyphema x 5 days
 - Hyphema >50% x 8 days

What complication is likely to occur if the IOP is >25 for longer than 5 days?

Corneal bloodstaining





Q

Hyphema

- *Hyphema: AC washout*

- Goal is to remove **circulating RBCs** from AC, not the **entire clot**
- Criteria triggering AC washout:
 - Any sign of **corneal bloodstaining**
 - Total hyphema x **5 days**
 - Hyphema >50% x **8 days**
 - Average IOP greater than:
 - 60 x **amount of time**



A

Hyphema

- *Hyphema: AC washout*
 - Goal is to remove **circulating RBCs** from AC, not the **entire clot**
 - Criteria triggering AC washout:
 - Any sign of **corneal bloodstaining**
 - Total hyphema x **5 days**
 - Hyphema >50% x **8 days**
 - Average IOP greater than:
 - 60 x **2 days**

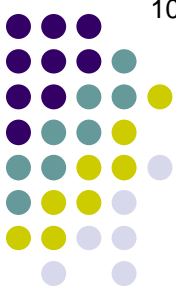


Q

Hyphema

- *Hyphema: AC washout*

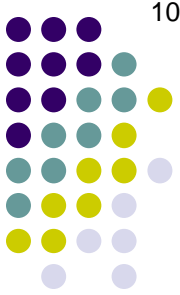
- Goal is to remove **circulating RBCs** from AC, not the **entire clot**
- Criteria triggering AC washout:
 - Any sign of **corneal bloodstaining**
 - Total hyphema x **5 days**
 - Hyphema >50% x **8 days**
 - Average IOP greater than:
 - 60 x **2 days** , or
 - 35 x **amount of time**



A

Hyphema

- *Hyphema: AC washout*
 - Goal is to remove **circulating RBCs** from AC, not the **entire clot**
 - Criteria triggering AC washout:
 - Any sign of **corneal bloodstaining**
 - Total hyphema x **5 days**
 - Hyphema >50% x **8 days**
 - Average IOP greater than:
 - 60 x **2 days** , or
 - 35 x **7 days**



Q

Hyphema

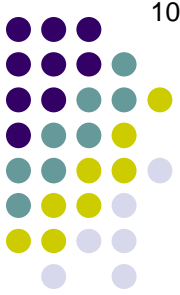
- *Hyphema: AC washout*

- Goal is to remove circulating RBCs from AC, not the entire clot
- Criteria triggering AC washout:
 - Any sign of corneal bloodstaining
 - Total hyphema x 5 days
 - Hyphema >50% x 8 days
 - Average IOP greater than:
 - 60 x 2 days , or
 - 35 x 7 days

What devastating complication is likely to occur if IOP is >60 for a couple of days, or >35 for seven days?

A

Hyphema



- *Hyphema: AC washout*

- Goal is to remove circulating RBCs from AC, not the entire clot
- Criteria triggering AC washout:
 - Any sign of corneal bloodstaining
 - Total hyphema x 5 days
 - Hyphema >50% x 8 days
 - Average IOP greater than:
 - 60 x 2 days , or
 - 35 x 7 days

What devastating complication is likely to occur if IOP is >60 for a couple of days, or >35 for seven days?
Optic atrophy

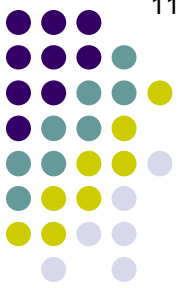


Q

Hyphema

- *Hyphema: AC washout*

- Goal is to remove **circulating RBCs** from AC, not the **entire clot**
- Criteria triggering AC washout:
 - Any sign of **corneal bloodstaining**
 - Total hyphema x **5 days**
 - Hyphema >50% x **8 days**
 - Average IOP greater than:
 - 60 x **2 days** , or
 - 35 x **7 days** , or
 - 25 x **amount of time**



A

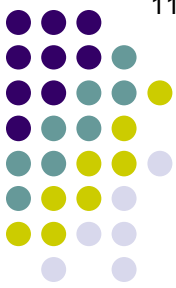
Hyphema

- *Hyphema: AC washout*
 - Goal is to remove **circulating RBCs** from AC, not the **entire clot**
 - Criteria triggering AC washout:
 - Any sign of **corneal bloodstaining**
 - Total hyphema x **5 days**
 - Hyphema >50% x **8 days**
 - Average IOP greater than:
 - 60 x **2 days** , or
 - 35 x **7 days** , or
 - 25 x **5 days**

Q

Hyphema

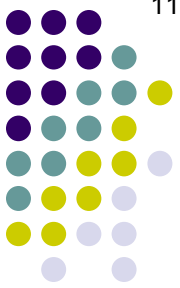
*What systemic condition **must** be checked for in hyphema pts at risk for it?*



A

Hyphema

*What systemic condition **must** be checked for in hyphema pts at risk for it?*
Sickle-cell anemia

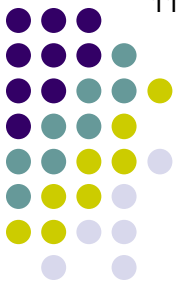


Q

Hyphema

*What systemic condition **must** be checked for in hyphema pts at risk for it?*
Sickle-cell anemia

Broadly speaking, what sort of disease is sickle-cell?

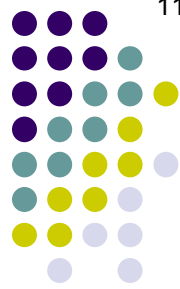


A

Hyphema

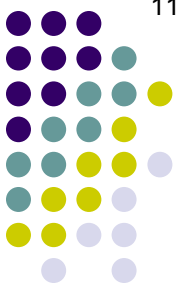
*What systemic condition **must** be checked for in hyphema pts at risk for it?*
Sickle-cell anemia

Broadly speaking, what sort of disease is sickle-cell?
A hemoglobinopathy



Q

Hyphema



*What systemic condition **must** be checked for in hyphema pts at risk for it?*
Sickle-cell anemia

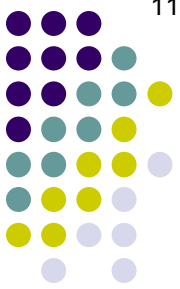
Broadly speaking, what sort of disease is sickle-cell?
A hemoglobinopathy

What is the underlying problem?

A

Hyphema

117

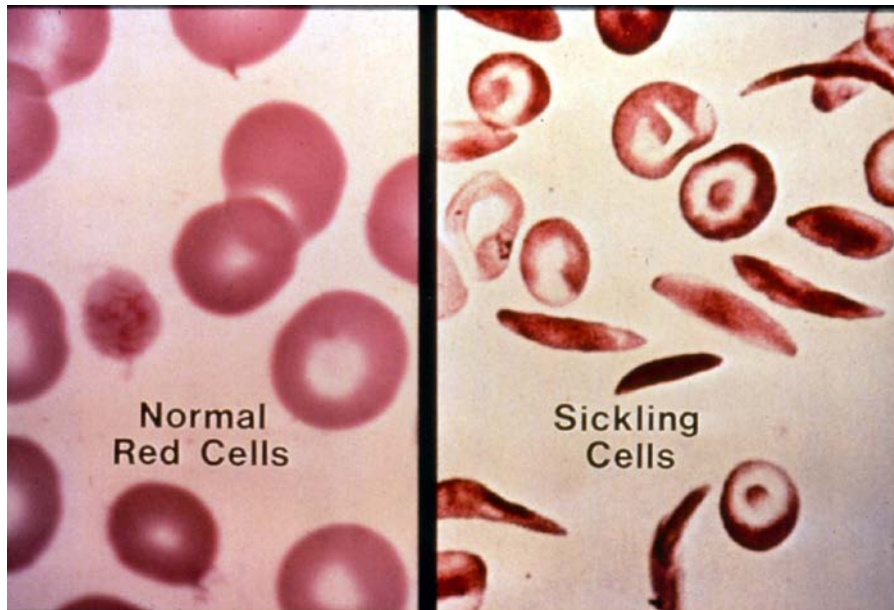


*What systemic condition **must** be checked for in hyphema pts at risk for it?*
Sickle-cell anemia

Broadly speaking, what sort of disease is sickle-cell?
A hemoglobinopathy

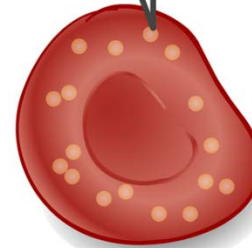
What is the underlying problem?
An amino-acid substitution in the hemoglobin beta-chain leads to its misfolding under certain metabolic conditions (eg, low O₂ tension). This results in the characteristic 'sickling' of affected RBCs.

Hyphema



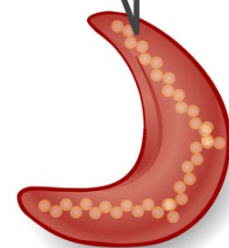
NORMAL RBC

Normal
hemoglobin
molecules



SICKLED RBC

Abnormal
hemoglobin
molecules

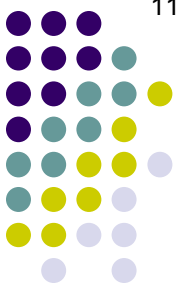


© AboutKidsHealth.ca

Sickle cell: RBC sickling



Hyphema



*What systemic condition **must** be checked for in hyphema pts at risk for it?*
Sickle-cell anemia

Broadly speaking, what sort of disease is sickle-cell?
A hemoglobinopathy

What is the underlying problem?
An amino-acid substitution in the hemoglobin beta-chain leads to its misfolding under certain metabolic conditions (eg, low O₂ tension). This results in the characteristic 'sickling' of affected RBCs.

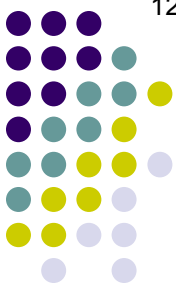
What are the four common genotypes of sickle-cell disease?

--
--
--
--

A

Hyphema

120



*What systemic condition **must** be checked for in hyphema pts at risk for it?*
Sickle-cell anemia

Broadly speaking, what sort of disease is sickle-cell?
A hemoglobinopathy

What is the underlying problem?
An amino-acid substitution in the hemoglobin beta-chain leads to its misfolding under certain metabolic conditions (eg, low O₂ tension). This results in the characteristic 'sickling' of affected RBCs.

What are the four common genotypes of sickle-cell disease?

- SS
- SC
- S-Thal
- SA

Q

Hyphema



What systemic condition **must** be checked for in hyphema pts at risk for it?
Sickle-cell anemia

Broadly speaking, what sort of disease is sickle-cell?
A hemoglobinopathy

What is the underlying problem?
An amino-acid substitution in the hemoglobin beta-chain leads to its malfolding under certain metabolic conditions (eg, low O₂ tension). This results in the characteristic 'sickling' of affected RBCs.

What are the four common genotypes of sickle-cell disease?

--SS
--SC
--S-Thal
--SA

What is the key difference between SS, SC and S-Thal vs SA disease?

A

Hyphema



What systemic condition **must** be checked for in hyphema pts at risk for it?
Sickle-cell anemia

Broadly speaking, what sort of disease is sickle-cell?
A hemoglobinopathy

What is the underlying problem?
An amino-acid substitution in the hemoglobin beta-chain leads to its misfolding under certain metabolic conditions (eg, low O₂ tension). This results in the characteristic 'sickling' of affected RBCs.

What are the four common genotypes of sickle-cell disease?

--SS
--SC
--S-Thal
--SA



What is the key difference between SS, SC and S-Thal vs SA disease?
The first three manifest as clinically apparent dz, whereas SA is an asymptomatic (under most conditions) carrier state--aka 'sickle trait'

Q

Hyphema



What systemic condition **must** be checked for in hyphema **pts at risk for it?**
Sickle-cell anemia

Broadly speaking, where? *In America, people of which two ethnic identities are at greatest risk?*
A hemoglobinopathy --
--

What is the underlying problem?

An amino-acid substitution in the hemoglobin beta-chain leads to its malfolding under certain metabolic conditions (eg, low O₂ tension). This results in the characteristic 'sickling' of affected RBCs.

What are the four common genotypes of sickle-cell disease?

- SS
- SC
- S-Thal
- SA

A

Hyphema



What systemic condition **must** be checked for in hyphema **pts at risk for it?**
Sickle-cell anemia

Broadly speaking, where? *In America, people of which two ethnic identities are at greatest risk?*
A hemoglobinopathy --African-American (People of Mediterranean and Southeast Asian ancestry
--Hispanic-American are also at some risk)

What is the underlying problem?

An amino-acid substitution in the hemoglobin beta-chain leads to its malfolding under certain metabolic conditions (eg, low O₂ tension). This results in the characteristic 'sickling' of affected RBCs.

What are the four common genotypes of sickle-cell disease?

- SS
- SC
- S-Thal
- SA

Q

Hyphema



What systemic condition **must** be checked for in hyphema **pts at risk for it?**
Sickle-cell anemia

Broadly speaking, where? In America, people of which two ethnic identities are at greatest risk?
A hemoglobinopathy --African-American: 1 in ?
--Hispanic-American: 1 in ?

What is the underlying mechanism? What is the sickle-cell dz birthrate for these groups?
An amino-acid substitution in the hemoglobin beta-chain leads to its misfolding under certain metabolic conditions (eg, low O₂ tension). This results in the characteristic 'sickling' of affected RBCs.

What are the four common genotypes of sickle-cell disease?

- SS
- SC
- S-Thal
- SA

A

Hyphema



What systemic condition **must** be checked for in hyphema *pts at risk for it?*
Sickle-cell anemia

Broadly speaking, where? In America, people of which two ethnic identities are at greatest risk?
A hemoglobinopathy --African-American: 1 in 500
--Hispanic-American: 1 in 36,000

What is the underlying mechanism? What is the sickle-cell dz birthrate for these groups?
An amino-acid substitution in the hemoglobin beta-chain leads to its misfolding under certain metabolic conditions (eg, low O₂ tension). This results in the characteristic 'sickling' of affected RBCs.

What are the four common genotypes of sickle-cell disease?

- SS
- SC
- S-Thal
- SA

Q

Hyphema



What systemic condition **must** be checked for in hyphema **pts at risk for it?**
Sickle-cell anemia

Broadly speaking, where? In America, people of which two ethnic identities are at greatest risk?
A hemoglobinopathy --African-American: 1 in 500
--Hispanic-American: 1 in 36,000

What is the underlying mechanism? What is the sickle-cell dz birthrate for these groups?
An amino-acid substitution in the hemoglobin beta-chain leads to its misfolding under certain metabolic conditions (eg, low O₂ tension). This results in the characteristic 'sickling' of affected RBCs.

What are the four common genotypes of sickle-cell disease?

--SS

--SC

--S-Thal

SA

What percent of African-Americans test positive for sickle trait?

A

Hyphema



What systemic condition **must** be checked for in hyphema **pts at risk for it?**
Sickle-cell anemia

Broadly speaking, where? In America, people of which two ethnic identities are at greatest risk?
A hemoglobinopathy --African-American: 1 in 500
--Hispanic-American: 1 in 36,000

What is the underlying mechanism? What is the sickle-cell dz birthrate for these groups?
An amino-acid substitution in the hemoglobin beta-chain leads to its misfolding under certain metabolic conditions (eg, low O₂ tension). This results in the characteristic 'sickling' of affected RBCs.

What are the four common genotypes of sickle-cell disease?

--SS

--SC

--S-Thal

--SA

What percent of African-Americans test positive for sickle trait?

A whopping 8% (1 in 12)!



Hyphema



*What systemic condition **must** be checked for in hyphema pts at risk for it?*
Sickle-cell anemia

Broadly speaking, what sort of disease is sickle-cell?
A hemoglobinopathy

What is the underlying problem?
An amino-acid substitution in the hemoglobin beta-chain leads to its malfunctioning under certain metabolic conditions (eg, low O₂ tension). This results in the characteristic 'sickling' of affected RBCs.

What are the four common genotypes of sickle-cell disease?

--SS

--SC

--S-Thal

--SA

Why must sickle-status be assessed in at-risk hyphema pts?

Because being sickle-positive:

1)

2)



*What systemic condition **must** be checked for in hyphema pts at risk for it?*
Sickle-cell anemia

Broadly speaking, what sort of disease is sickle-cell?
A hemoglobinopathy

What is the underlying problem?
An amino-acid substitution in the hemoglobin beta-chain leads to its misfolding under certain metabolic conditions (eg, low O₂ tension). This results in the characteristic 'sickling' of affected RBCs.

What are the four common genotypes of sickle-cell disease?
--SS
--SC
--S-Thal
--SA

Why must sickle-status be assessed in at-risk hyphema pts?
Because being sickle-positive:
1) places a hyphema pt at higher risk for complications; and
2) impacts how a hyphema should be managed

Q

Hyphema



What systemic condition **must** be checked for in hyphema pts at risk for it?
Sickle-cell anemia

Broadly speaking, what sort of disease is sickle-cell?
A hemoglobinopathy

What is the underlying problem?
An amino-acid substitution in the hemoglobin beta-chain leads to its malfolding under certain metabolic conditions (eg, low O₂ tension). This results in the characteristic 'sickling' of affected RBCs.

What are the four common genotypes of sickle-cell disease?

--SS?
--SC?
--S-Thal?
--SA?

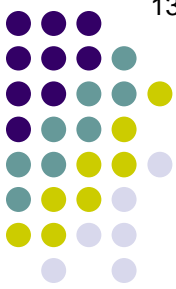
Which is/are associated with increased risk and a need for modified management?

Why must sickle-status be assessed in at-risk hyphema pts?
Because being sickle-positive:

1) places a hyphema pt at higher risk for complications; and
2) impacts how a hyphema should be managed

A

Hyphema



What systemic condition **must** be checked for in hyphema pts at risk for it?
Sickle-cell anemia

Broadly speaking, what sort of disease is sickle-cell?
A hemoglobinopathy

What is the underlying problem?
An amino-acid substitution in the hemoglobin beta-chain leads to its malfolding under certain metabolic conditions (eg, low O₂ tension). This results in the characteristic 'sickling' of affected RBCs.

What are the four common genotypes of sickle-cell disease?

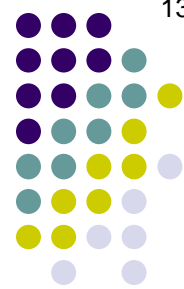
--SS!
--SC!
--S-Thal!
--SA!

Which is/are associated with increased risk and a need for modified management?

All of them

Why must sickle-status be assessed in at-risk hyphema pts?
Because being sickle-positive:

1) places a hyphema pt at higher risk for complications; and
2) impacts how a hyphema should be managed



Q

Hyphema

What systemic conditions are sickle-cell anemia pts at risk for it?
Sickle-cell anemia

What two complications are sickle pts at greater risk of developing?

Broadly speaking, what sort of disease is sickle-cell?
A hemoglobinopathy

What is the underlying problem?

An amino-acid substitution in the hemoglobin beta-chain leads to its malfunctioning under certain metabolic conditions (eg, low O₂ tension). This results in the characteristic 'sickling' of affected RBCs.

What are the four common genotypes of sickle-cell disease?

--SS!
--SC!
--S-Thal!
--SA!

Why must sickle-status be assessed in at-risk hyphema pts?

Because being sickle-positive:

1) places a hyphema pt at higher risk for complications; and
2) impacts how a hyphema should be managed



A

Hyphema

What systemic conditions are sickle-cell pts at risk for it?
Sickle-cell anemia

What two complications are sickle pts at greater risk of developing?
Elevated IOP and optic atrophy

Broadly speaking, what sort of disease is sickle-cell?
A hemoglobinopathy

What is the underlying problem?
An amino-acid substitution in the hemoglobin beta-chain leads to its malfunctioning under certain metabolic conditions (eg, low O₂ tension). This results in the characteristic 'sickling' of affected RBCs.

What are the four common genotypes of sickle-cell disease?
--SS!
--SC!
--S-Thal!
--SA!

Why must sickle-status be assessed in at-risk hyphema pts?
Because being sickle-positive:

1) places a hyphema pt at higher risk for complications; and
2) impacts how a hyphema should be managed



Q

Hyphema

What systemic conditions are sickle-cell anemia pts at risk for it?
Sickle-cell anemia

Elevated IOP and optic atrophy

Broadly speaking, what sort of disease is sickle-cell?

Why do sickle pts have an increased risk of developing elevated IOP?

What are the four common genotypes of sickle-cell disease?

- SS!
- SC!
- S-Thal!
- SA!

Why must sickle-status be assessed in at-risk hyphema pts?

Because being sickle-positive:

- 1) places a hyphema pt at higher risk for complications; and
- 2) impacts how a hyphema should be managed



A

Hyphema

What systemic conditions are sickle-cell anemia pts at risk for it?
Sickle-cell anemia

Elevated IOP and optic atrophy

Broadly speaking, what sort of disease is sickle-cell?

Why do sickle pts have an increased risk of developing elevated IOP?

Because of the physical characteristics of their RBCs. Normal RBCs are very flexible and change shape easily; thus, they are able to slip easily through the TM.

What are the four common genotypes of sickle-cell disease?

- SS!
- SC!
- S-Thal!
- SA!

Why must sickle-status be assessed in at-risk hyphema pts?

Because being sickle-positive:

- 1) places a hyphema pt at higher risk for complications;** and
- 2) impacts how a hyphema should be managed**



A

Hyphema

What systemic conditions are sickle-cell pts at risk for it?
Sickle-cell anemia

Elevated IOP and optic atrophy

Broadly speaking, what sort of disease is sickle-cell?

Why do sickle pts have an increased risk of developing elevated IOP?

Because of the physical characteristics of their RBCs. Normal RBCs are very flexible and change shape easily; thus, they are able to slip easily through the TM. In contrast, sickled RBCs are stiff, and therefore cannot readily change conformation to fit through the TM. Thus, sickled RBCs will 'pile up' at the TM, effectively occluding it and thereby causing IOP to spike.

What are the four common genotypes of sickle-cell disease?

- SS!
- SC!
- S-Thal!
- SA!

Why must sickle-status be assessed in at-risk hyphema pts?

Because being sickle-positive:

- 1) places a hyphema pt at higher risk for complications;** and
- 2) impacts how a hyphema should be managed**



Q

Hyphema

What systemic conditions are pts at risk for it?
Sickle-cell anemia

Elevated IOP and **optic atrophy**

Broadly speaking, what sort of disease is sickle-cell?

Why do sickle pts have an increased risk of developing elevated IOP?

Because of the physical characteristics of their RBCs. Normal RBCs are very flexible and change shape easily; thus, they are able to slip easily through the TM. In contrast, sickled RBCs are stiff, and therefore cannot readily change conformation to fit through the TM. Thus, sickled RBCs will 'pile up' at the TM, effectively occluding it and thereby causing IOP to spike.

What are the four complications of sickle-cell anemia?
Why are the optic nerves in sickle pts at greater risk of atrophying?

--SS!
--SC!
--S-Thal!
--SA!

Why must sickle-state be managed?
Because being sickle positive.

1) places a hyphema pt at higher risk for complications; and
2) impacts how a hyphema should be managed

A

Hyphema



What systemic conditions are sickle-cell pts at risk for it?
Sickle-cell anemia

Elevated IOP and **optic atrophy**

Broadly speaking, what sort of disease is sickle-cell?

Why do sickle pts have an increased risk of developing elevated IOP?
Because of the physical characteristics of their RBCs. Normal RBCs are very flexible and change shape easily; thus, they are able to slip easily through the TM. In contrast, sickled RBCs are stiff, and therefore cannot readily change conformation to fit through the TM. Thus, sickled RBCs will 'pile up' at the TM, effectively occluding it and thereby causing IOP to spike.

What are the four complications?

- SS!
- SC!
- S-Thal!
- SA!

Why must sickle-state be managed?

Because being sickle positive.

Why are the optic nerves in sickle pts at greater risk of atrophying?

As is the case throughout their bodies, the baseline status of the microcirculation of the optic nerve in sicklers is marginal—even under the best of circumstances, their optic nerves are just getting by. Thus, their optic nerves are vulnerable to damage stemming from anything that compromises their already compromised circulation, one of which is elevated IOP.

- 1) places a hyphema pt at higher risk for complications; and
- 2) impacts how a hyphema should be managed

A

Hyphema



What systemic conditions are sickle-cell pts at risk for it?
Sickle-cell anemia

Elevated IOP

optic atrophy

Broadly speaking, what sort of disease is sickle-cell?

Why do sickle pts have an increased risk of developing elevated IOP?

Because of the physical characteristics of their RBCs. Normal RBCs are very flexible and change shape easily; thus, they are able to slip easily through the TM. In contrast, sickled RBCs are stiff, and therefore cannot readily change conformation to fit through the TM. Thus, sickled RBCs will 'pile up' at the TM, effectively occluding it and thereby causing IOP to spike.

What are the four complications?

- SS!
- SC!
- S-Thal!
- SA!

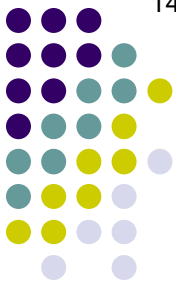
Why must sickle-state be managed?

Because being sickle-positive.

Why are the optic nerves in sickle pts at greater risk of atrophying?

As is the case throughout their bodies, the baseline status of the microcirculation of the optic nerve in sicklers is marginal—even under the best of circumstances, their optic nerves are just getting by. Thus, their optic nerves are vulnerable to damage stemming from anything that compromises their already compromised circulation, one of which is elevated IOP. Because of this, the optic nerve in sickle pts will sustain **more** damage at **lower** IOPs for **shorter** durations than will the optic nerves of non-sickle pts.

- 1) places a hyphema pt at higher risk for complications; and
- 2) impacts how a hyphema should be managed



Q

Hyphema

What systemic condition **must** be checked for in hyphema pts at risk for it?
Sickle-cell anemia

How does being sickle+ impact hyphema management?

?

What is the underlying problem?

An amino-acid substitution in the hemoglobin beta-chain leads to its malfolding under certain metabolic conditions (eg, low O₂ tension). This results in the characteristic 'sickling' of affected RBCs.

What are the four common genotypes of sickle-cell disease?

--SS!

--SC!

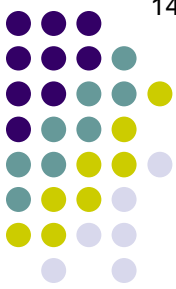
--S-Thal!

--SA!

Why must sickle-status be assessed in at-risk hyphema pts?

Because being sickle-positive:

- 1) places a hyphema pt at higher risk for complications; and
- 2) impacts how a hyphema should be managed



A

Hyphema

What systemic condition **must** be checked for in hyphema pts at risk for it?
Sickle-cell anemia

How does being sickle+ impact hyphema management?

By changing:

- The agents employed in *medical* management, and
- The threshold for advancing to *surgical* management

?

What is the underlying problem?

An amino-acid substitution in the hemoglobin beta-chain leads to its malfolding under certain metabolic conditions (eg, low O₂ tension). This results in the characteristic 'sickling' of affected RBCs.

What are the four common genotypes of sickle-cell disease?

- SS!
- SC!
- S-Thal!
- SA!

Why must sickle-status be assessed in at-risk hyphema pts?

Because being sickle-positive:

- 1) places a hyphema pt at higher risk for complications; and
- 2) impacts how a hyphema should be managed

Q

Hyphema

What systemic condition **must** be checked for in hyphema pts at risk for it?
Sickle-cell anemia

How does being sickle+ impact hyphema management?

By changing:

- The agents employed in *medical* management and
- The threshold for advancing to *surgical* management

Which two classes of meds are first-line in managing IOP in hyphema?

These are the agents commonly employed in managing IOP in hyphema.
Which ones should be **avoided** in sicklers?

--Aqueous suppressants

---- β blockers?

---- α_2 agonists

-----Apraclonidine (iopidine)?

-----Brimonidine?

----Carbonic anhydrase inhibitors (CAIs)?

--Hyperosmotic agents?

----Mannitol?

----Glycerol?

1) places a hyphema pt at higher risk for complications; and

2) impacts how a hyphema should be managed

A

Hyphema



What systemic condition **must** be checked for in hyphema pts at risk for it?

Sickle-cell anemia

How does being sickle+ impact hyphema management?

By changing:

- The agents employed in *medical* management and
- The threshold for advancing to *surgical* management

Which two classes of meds are first-line in managing IOP in hyphema?

These are the agents commonly employed in managing IOP in hyphema.
Which ones should be **avoided** in sicklers?

--Aqueous suppressants

---- β blockers

---- α_2 agonists

-----Apraclonidine (iopidine)!

-----Brimonidine

----Carbonic anhydrase inhibitors (CAIs)!

--Hyperosmotic agents?

----Mannitol!

----Glycerol!

Remember: These are the agents **not** to use in sicklers!

1) places a hyphema pt at higher risk for complications; and

2) impacts how a hyphema should be managed



Q

Hyphema

What systemic condition **must** be checked for in hyphema pts at risk for it?

Sickle-cell anemia

How does being sickle+ impact hyphema management?

By changing.

- The agents employed in medical management, and
- The threshold for advancing to surgical management

Which two classes of meds are first-line in managing IOP in hyphema?

These are the agents commonly employed in managing IOP in hyphema.
Which ones should be avoided in sicklers?

--Aqueous suppressants

---- β blockers

---- α_1 agonists

----Apraclonidine (iopidine)

----Brimonidine

----Carbonic anhydrase inhibitors

--Hyperosmotic agents?

----Mannitol

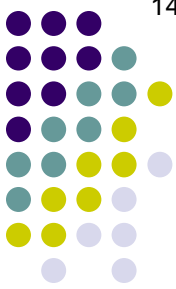
----Glycerol

Why must apraclonidine be avoided in sicklers?

- 1) places a hyphema pt at higher risk for complications; and
- 2) impacts how a hyphema should be managed

Q/A

Hyphema



What systemic condition **must** be checked for in hyphema pts at risk for it?
Sickle-cell anemia

How does being sickle+ impact hyphema management?

By changing.

- The agents employed in medical management, and
- The threshold for advancing to surgical management

Which two classes of meds are first-line in managing IOP in hyphema?

These are the agents commonly employed in managing IOP in hyphema.
Which ones should be avoided in sicklers?

--Aqueous suppressants

---- β blockers

---- α_1 agonists

----Apraclonidine (iopidine)

----Brimonidine

----Carbonic anhydrase inhibitors

--Hyperosmotic agents?

----Mannitol

----Glycerol

Why must apraclonidine be avoided in sicklers?

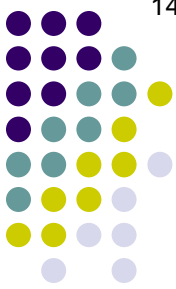
It has too much stimulatory effect

receptor class

- 1) places a hyphema pt at higher risk for complications; and
- 2) impacts how a hyphema should be managed

A

Hyphema



What systemic condition **must** be checked for in hyphema pts at risk for it?

Sickle-cell anemia

How does being sickle+ impact hyphema management?

By changing.

- The agents employed in medical management, and
- The threshold for advancing to surgical management

Which two classes of meds are first-line in managing IOP in hyphema?

These are the agents commonly employed in managing IOP in hyphema.
Which ones should be avoided in sicklers?

--Aqueous suppressants

---- β blockers

---- α_1 agonists

----Apraclonidine (iopidine)

-----Brimonidine

----Carbonic anhydrase inhibitors

--Hyperosmotic agents?

----Mannitol

----Glycerol

Why must apraclonidine be avoided in sicklers?

It has too much α_1 stimulatory effect

- 1) places a hyphema pt at higher risk for complications; and
- 2) impacts how a hyphema should be managed

Q

Hyphema



What systemic condition **must** be checked for in hyphema pts at risk for it?
Sickle-cell anemia

How does being sickle+ impact hyphema management?

By changing.

- The agents employed in medical management, and
- The threshold for advancing to surgical management

Which two classes of meds are first-line in managing IOP in hyphema?

These are the agents commonly employed in managing IOP in hyphema.
Which ones should be avoided in sicklers?

--Aqueous suppressants

---- β blockers

---- α_1 agonists

----Apraclonidine (iopidine)

-----Brimonidine

----Carbonic anhydrase inhibitors

--Hyperosmotic agents?

----Mannitol

----Glycerol

Why must apraclonidine be avoided in sicklers?

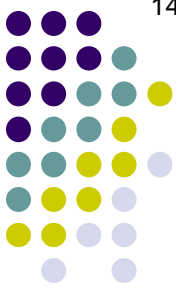
It has too much α_1 stimulatory effect

Which α_1 stimulation effect is so deleterious that it precludes the use of apraclonidine in sicklers?

- 1) places a hyphema pt at higher risk for complications; and
- 2) impacts how a hyphema should be managed

A

Hyphema



What systemic condition **must** be checked for in hyphema pts at risk for it?

Sickle-cell anemia

How does being sickle+ impact hyphema management?

By changing.

- The agents employed in medical management, and
- The threshold for advancing to surgical management

Which two classes of meds are first-line in managing IOP in hyphema?

These are the agents commonly employed in managing IOP in hyphema.
Which ones should be avoided in sicklers?

--Aqueous suppressants

---- β blockers

---- α_1 agonists

----Apraclonidine (iopidine)

-----Brimonidine

----Carbonic anhydrase inhibitors

--Hyperosmotic agents?

----Mannitol

----Glycerol

Why must apraclonidine be avoided in sicklers?

It has too much α_1 stimulatory effect

Which α_1 stimulation effect is so deleterious that it precludes the use of apraclonidine in sicklers?

Vasoconstriction of the anterior segment vasculature

- 1) places a hyphema pt at higher risk for complications; and
- 2) impacts how a hyphema should be managed

Q

Hyphema



What systemic condition **must** be checked for in hyphema pts at risk for it?
Sickle-cell anemia

How does being sickle+ impact hyphema management?

By changing.

- The agents employed in medical management, and
- The threshold for advancing to surgical management

Which two classes of meds are first-line in managing IOP in hyphema?

These are the agents commonly employed in managing IOP in hyphema.
Which ones should be **avoided** in sicklers?

--Aqueous suppressants

---- β blockers

---- α_2 agonists

-----Apraclonidine (iopidine)

-----Brimonidine

-----**Carbonic anhydrase inhibitors (CAIs)**

--Hyperosmotic agents?

----Mannitol

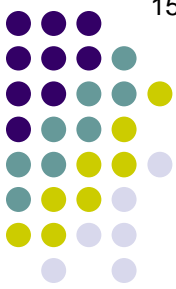
----Glycerol

Why should topical CAIs be avoided in sicklers?

- 1) places a hyphema pt at higher risk for
- 2) impacts how a hyphema should be managed

A

Hyphema



What systemic condition **must** be checked for in hyphema pts at risk for it?
Sickle-cell anemia

How does being sickle+ impact hyphema management?

By changing.

- The agents employed in medical management, and
- The threshold for advancing to surgical management

Which two classes of meds are first-line in managing IOP in hyphema?

These are the agents commonly employed in managing IOP in hyphema.
Which ones should be **avoided** in sicklers?

--Aqueous suppressants

---- β blockers

---- α_2 agonists

-----Apraclonidine (iopidine)

-----Brimonidine

-----**Carbonic anhydrase inhibitors (CAIs)**

--Hyperosmotic agents?

----Mannitol

----Glycerol

Why should topical CAIs be avoided in sicklers?

Because they will acidify the AC, and an acidic environment promotes RBC sickling

- 1) places a hyphema pt at higher risk for
- 2) impacts how a hyphema should be managed

Q

Hyphema



What systemic condition **must** be checked for in hyphema pts at risk for it?
Sickle-cell anemia

How does being sickle+ impact hyphema management?

By changing.

- The agents employed in medical management, and
- The threshold for advancing to surgical management

Which two classes of meds are first-line in managing IOP in hyphema?

These are the agents commonly employed in managing IOP in hyphema.
Which ones should be **avoided** in sicklers?

--Aqueous suppressants

---- β blockers

---- α_2 agonists

-----Apraclonidine (iopidine)

-----Brimonidine

-----**Carbonic anhydrase inhibitors (CAIs)**

--Hyperosmotic agents?

----Mannitol

----Glycerol

Why should topical CAIs be avoided in sicklers?

Because they will acidify the AC, and an acidic environment promotes RBC sickling

There's an additional reason **systemic** CAIs should be avoided—what is it?

1) places a hyphema pt at higher risk for

2) impacts how a hyphema should be managed

A

Hyphema



What systemic condition **must** be checked for in hyphema pts at risk for it?
Sickle-cell anemia

How does being sickle+ impact hyphema management?

By changing.

- The agents employed in medical management, and
- The threshold for advancing to surgical management

Which two classes of meds are first-line in managing IOP in hyphema?

These are the agents commonly employed in managing IOP in hyphema.
Which ones should be **avoided** in sicklers?

--Aqueous suppressants

---- β blockers

---- α_2 agonists

-----Apraclonidine (iopidine)

-----Brimonidine

-----**Carbonic anhydrase inhibitors (CAIs)**

--Hyperosmotic agents?

----Mannitol

----Glycerol

Why should topical CAIs be avoided in sicklers?

Because they will acidify the AC, and an acidic environment promotes RBC sickling

There's an additional reason **systemic** CAIs should be avoided—what is it?

Because of their diuretic effect, they may lead to hemoconcentration and thus circulatory compromise (especially if the pt is already dehydrated)

- 1) places a hyphema pt at higher risk for
- 2) impacts how a hyphema should be managed



Q

Hyphema

What systemic condition **must** be checked for in hyphema pts at risk for it?

Sickle-cell anemia

How does being sickle+ impact hyphema management?

By changing.

- The agents employed in medical management, and
- The threshold for advancing to surgical management

Which two classes of meds are first-line in managing IOP in hyphema?

These are the agents commonly employed in managing IOP in hyphema. Which ones should be **avoided** in sicklers?

- Aqueous suppressants
- β blockers
- α_2 agonists
- Apraclonidine (iopidine)
- Brimonidine
- Carbonic anhydrase inhibitors (CAIs)
- Hyperosmotic agents?

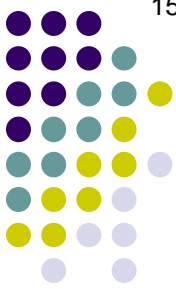
Mannitol
Glycerol

Why should the hyperosmotics be avoided?

- 1) places a hyphema pt at higher risk for complications, and
- 2) impacts how a hyphema should be managed

A

Hyphema



What systemic condition **must** be checked for in hyphema pts at risk for it?

Sickle-cell anemia

How does being sickle+ impact hyphema management?

By changing.

- The agents employed in medical management, and
- The threshold for advancing to surgical management

Which two classes of meds are first-line in managing IOP in hyphema?

These are the agents commonly employed in managing IOP in hyphema.
Which ones should be **avoided** in sicklers?

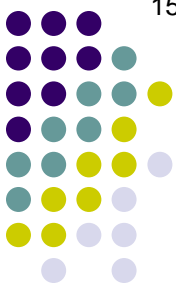
- Aqueous suppressants
- β blockers
- α_2 agonists
- Apraclonidine (iopidine)
- Brimonidine
- Carbonic anhydrase inhibitors (CAIs)
- Hyperosmotic agents?

Mannitol
Glycerol

Why should the hyperosmotics be avoided?

For the same reason as systemic CAIs--they may lead to hemoconcentration and thus circulatory compromise

- 1) places a hyphema pt at higher risk for complications, and
- 2) impacts how a hyphema should be managed



Hyphema

What systemic condition **must** be checked for in hyphema pts at risk for it?

Sickle-cell anemia

How does being sickle+ impact hyphema management?

By changing:

- The agents employed in medical management, and
- The threshold for advancing to surgical management**

What is the underlying problem?

An amino acid substitution in the hemoglobin beta chain leads to its malfunctioning (sickling). This results in the

Next let's take a look at the IOP thresholds for doing an AC washout in sickle-cell pts

What are the four common genotypes of sickle-cell disease?

- SS!
- SC!
- S-Thal!
- SA!

Why must sickle-status be assessed in at-risk hyphema pts?

Because being sickle-positive:

- 1) places a hyphema pt at higher risk for complications; and
- 2) **impacts how a hyphema should be managed**

(No question—proceed when ready)



Q

Hyphema

• *Hyphema: AC washout*

- Goal is to remove circulating RBCs from AC, not the entire clot
- Criteria triggering AC washout:
 - Any sign of corneal bloodstaining
 - Total hyphema x 5 days
 - Hyphema >50% x 8 days
 - Average IOP greater than:
 - 60 x 2 days, or
 - 35 x 7 days, or
 - 25 x 5 days



If sickle-positive: Average IOP greater than:



- 25 x amount of time



A

Hyphema

• *Hyphema: AC washout*

- Goal is to remove circulating RBCs from AC, not the entire clot
- Criteria triggering AC washout:
 - Any sign of corneal bloodstaining
 - Total hyphema x 5 days
 - Hyphema >50% x 8 days
 - Average IOP greater than:
 - 60 x 2 days, or
 - 35 x 7 days, or
 - 25 x 5 days

★ ***If sickle-positive:*** Average IOP greater than: ★

- 25 x 1 day



Q

Hyphema

• *Hyphema: AC washout*

- Goal is to remove circulating RBCs from AC, not the entire clot
- Criteria triggering AC washout:
 - Any sign of corneal bloodstaining
 - Total hyphema x 5 days
 - Hyphema >50% x 8 days
 - Average IOP greater than:
 - 60 x 2 days, or
 - 35 x 7 days, or
 - 25 x 5 days



If sickle-positive: Average IOP greater than:



- 25 x 1 day, or
- Repeated spikes >30 for amount of time



A

Hyphema

• *Hyphema: AC washout*

- Goal is to remove circulating RBCs from AC, not the entire clot
- Criteria triggering AC washout:
 - Any sign of corneal bloodstaining
 - Total hyphema x 5 days
 - Hyphema >50% x 8 days
 - Average IOP greater than:
 - 60 x 2 days, or
 - 35 x 7 days, or
 - 25 x 5 days

★ ***If sickle-positive:*** Average IOP greater than: ★

- 25 x 1 day, or
- Repeated spikes >30 for 2 days



Hyphema

● *Hyphema: AC washout*

- Goal is to remove circulating RBCs from AC, not the entire clot
- Criteria triggering AC washout:
 - Any sign of corneal bloodstaining
 - Total hyphema x 5 days
 - Hyphema >50% x 8 days
 - Average IOP greater than:

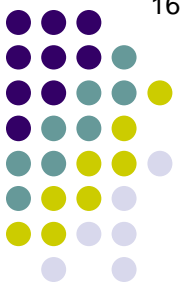
Note the management dilemma in sickle-cell—IOP control is absolutely essential because of their pre-existing optic-nerve vulnerability, but sickle-status takes most medical IOP-lowering agents out of your hands

- 25 x 5 days

★ **If sickle-positive: Average IOP greater than:** ★

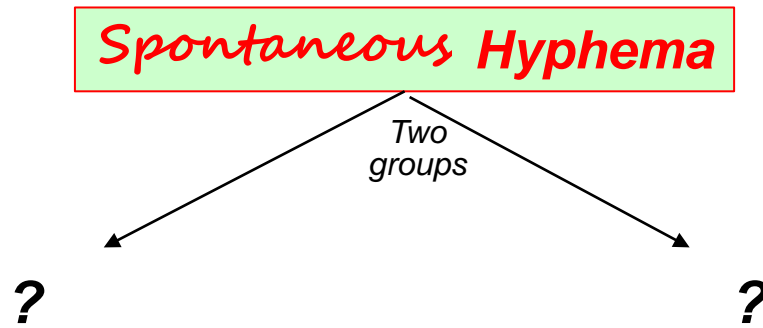
- 25 x 1 day , or
- Repeated spikes >30 for 2 days

Spontaneous Hyphema

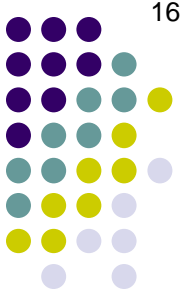


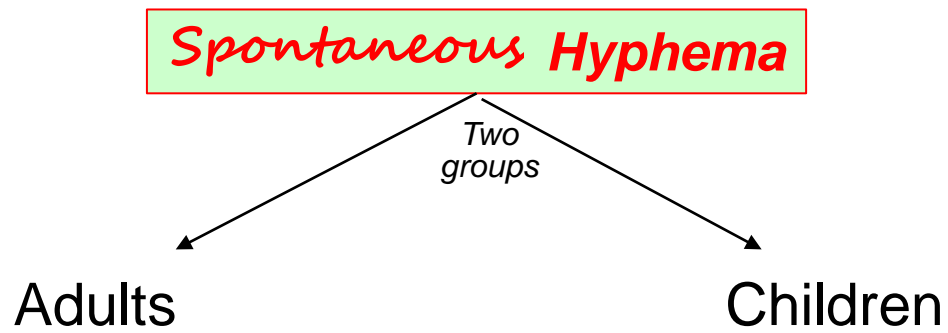
Next, let's turn our attention to the subject of
spontaneous hyphema

Q

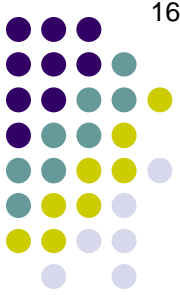


When considering the DDx for spontaneous hyphema, we need to think in terms of two groups. What are they?



A

When considering the DDx for spontaneous hyphema, we need to think in terms of two groups. What are they?



Q

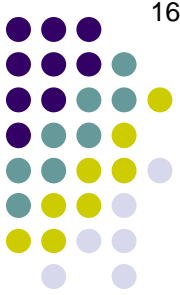
Spontaneous Hyphema

Adults

└ ?
└ ?
└ ?

Children

What are the three most common causes of spontaneous hyphema in adults?



A

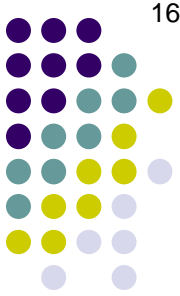
Spontaneous Hyphema

Adults

- └ Neovascularization
- └ Neovascularization
- └ Neovascularization

Children

What are the three most common causes of spontaneous hyphema in adults?





Q

Spontaneous Hyphema

Adults

- └ Neovascularization
- └ Neovascularization
- └ Neovascularization

Children

What are the three most common causes of spontaneous hyphema in adults?

Yo Dr Flynn, you listed neovascularization three times. This a mistake, or what?



A

Spontaneous Hyphema

Adults

- └ Neovascularization
- └ Neovascularization
- └ Neovascularization

Children

What are the three most common causes of spontaneous hyphema in adults?

Yo Dr Flynn, you listed neovascularization three times. This a mistake, or what?
It's 'or what,' actually. The point being made here is that neovascularization of the anterior segment is the final common pathway for the conditions that commonly produce spontaneous hyphema in adults.

Q

Spontaneous Hyphema

Adults

- └ Neovascularization 2ndry to...
- └ Neovascularization 2ndry to...
- └ Neovascularization 2ndry to...

Children

Yo Dr Flynn, you listed neovascularization three times. This a mistake, or what?
 It's 'or what,' actually. The point being made here is that neovascularization of the anterior segment is the final common pathway for the conditions that commonly produce spontaneous hyphema in adults.

OK, then, what are the three sorts of common conditions that produce the neo that produces spontaneous hyphema in adults?

--
 --
 --





A

Spontaneous Hyphema

Adults

Children

- └ Neovascularization 2ndry to...ischemia
- └ Neovascularization 2ndry to...neoplasm
- └ Neovascularization 2ndry to...inflammation

Yo Dr Flynn, you listed neovascularization three times. This a mistake, or what?
It's 'or what,' actually. The point being made here is that neovascularization of the anterior segment is the final common pathway for the conditions that commonly produce spontaneous hyphema in adults.

OK, then, what are the three sorts of common conditions that produce the neo that produces spontaneous hyphema in adults?

- Ischemia
- Neoplasm
- Inflammation

Q

Spontaneous Hyphema

Adults

- └ Ischemia
- └ Neoplasm
- └ Inflammation

Children

? ➡ HYPHEMA

Speaking of common pathways in hyphema...Let's work backwards through the process that leads to free blood in the AC.

What event sets up the eye to get a hyphema?



A

Spontaneous Hyphema

Adults

- └ Ischemia
- └ Neoplasm
- └ Inflammation

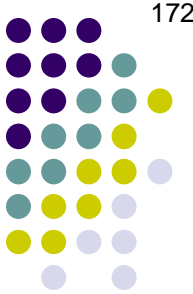
Children

A.S. Neo → HYPHEMA

Speaking of common pathways in hyphema...Let's work backwards through the process that leads to free blood in the AC.

What event sets up the eye to get a hyphema?

Neovascularization of the anterior segment



Q

Spontaneous Hyphema

Adults

- └ Ischemia
- └ Neoplasm
- └ Inflammation

Children

?

➡ A.S. Neo ➡ HYPHEMA

Speaking of common pathways in hyphema...Let's work backwards through the process that leads to free blood in the AC.

What event sets up the eye to get a hyphema?

Neovascularization of the anterior segment

What incites the development of anterior segment neo?



A

Spontaneous Hyphema

Adults

- └ Ischemia
- └ Neoplasm
- └ Inflammation

Children

Release of VEGF, other
vascular growth factors → A.S.
Neo → HYPHEMA

Speaking of common pathways in hyphema...Let's work backwards through the process that leads to free blood in the AC.

What event sets up the eye to get a hyphema?

Neovascularization of the anterior segment

What incites the development of anterior segment neo?

The presence of VEGF and other vascular growth factors



Q

Spontaneous Hyphema

Adults

- └ Ischemia
- └ Neoplasm
- └ Inflammation

Children

? ➡ Release of VEGF, other
vascular growth factors ➡ A.S.
Neo ➡ HYPHEMA

Speaking of common pathways in hyphema...Let's work backwards through the process that leads to free blood in the AC.

What event sets up the eye to get a hyphema?

Neovascularization of the anterior segment

What incites the development of anterior segment neo?

The presence of VEGF and other vascular growth factors

What event leads to the release of VEGF, etc?

A

Spontaneous Hyphema

Adults

- └ Ischemia
- └ Neoplasm
- └ Inflammation

Children

Hypoxia → Release of VEGF, other vascular growth factors → A.S. Neo → HYPHEMA

Speaking of common pathways in hyphema...Let's work backwards through the process that leads to free blood in the AC.

What event sets up the eye to get a hyphema?

Neovascularization of the anterior segment

What incites the development of anterior segment neo?

The presence of VEGF and other vascular growth factors

What event leads to the release of VEGF, etc?

Hypoxia



Q

Spontaneous Hyphema

Adults

Children

- Ischemia
- Neoplasm
- Inflammation

Hypoxia → Release of VEGF, other vascular growth factors → A.S. Neo → HYPHEMA

Broadly speaking, ischemia of one of two 'structures' (note the 'scare quotes') is responsible for producing anterior segment neo. What are these structures?



A

Spontaneous Hyphema

Adults

Children

└ Ischemia

└ Neoplasm

└ Inflammation

Hypoxia

Release of VEGF, other
vascular growth factors

A.S.
Neo

HYPHEMA

Broadly speaking, ischemia of one of two 'structures' (note the 'scare quotes') is responsible for producing anterior segment neo. What are these structures?

The Retina, and the Entire Eye



Q

Spontaneous Hyphema

Adults

Children

└ **Ischemia**

└ Neoplasm

└ Inflammation

Hypoxia → Release of VEGF, other
vascular growth factors → A.S.
Neo → HYPHEMA

*Broadly speaking, ischemia of one of two 'structures' (note the 'scare quotes') is responsible for producing anterior segment neo.
What are these structures?*

The **Retina**, and the Entire Eye

What retinal ischemic events are notorious for producing anterior segment neo?



A

Spontaneous Hyphema

Adults

Children

└ Ischemia

└ Neoplasm

└ Inflammation

Hypoxia → Release of VEGF, other vascular growth factors → A.S. Neo → HYPHEMA

Broadly speaking, ischemia of one of two 'structures' (note the 'scare quotes') is responsible for producing anterior segment neo. What are these structures?

The **Retina**, and the Entire Eye

What retinal ischemic events are notorious for producing anterior segment neo?

Proliferative diabetic retinopathy (PDR), and retinal vein occlusions—either central (CRVO) or branch (BRVO)



Q

Spontaneous Hyphema

Adults

Children

└ Ischemia

└ Neoplasm

└ Inflammation

Hypoxia

Release of VEGF, other
vascular growth factors

A.S.
Neo

HYPHEMA

Broadly speaking, ischemia of one of two 'structures' (note the 'scare quotes') is responsible for producing anterior segment neo. What are these structures?

The Retina, and the **Entire Eye**

What is the name of the condition in which the entire eye is ischemic?

A

Spontaneous Hyphema

Adults

Children

└ Ischemia

└ Neoplasm

└ Inflammation

Hypoxia

➡ Release of VEGF, other
vascular growth factors ➡ A.S.
Neo ➡ HYPHEMA

Broadly speaking, ischemia of one of two 'structures' (note the 'scare quotes') is responsible for producing anterior segment neo. What are these structures?

The Retina, and the **Entire Eye**

What is the name of the condition in which the entire eye is ischemic?
Ocular ischemic syndrome (OIS)

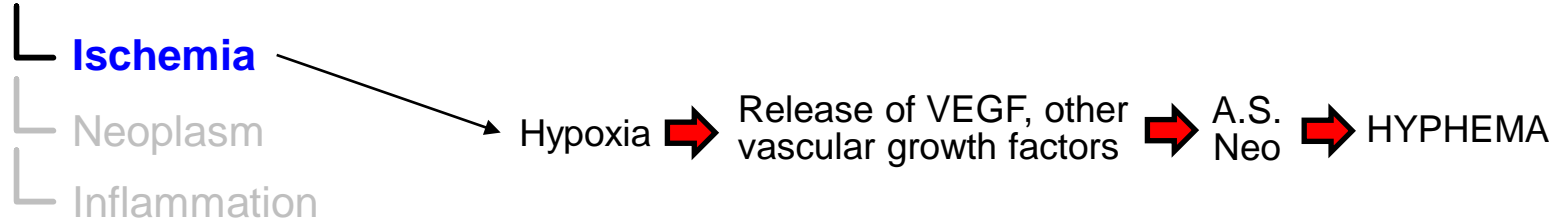


Q

Spontaneous Hyphema

Adults

Children



Broadly speaking, ischemia of one of two 'structures' (note the 'scare quotes') is responsible for producing anterior segment neo. What are these structures?

The Retina, and the **Entire Eye**

What is the name of the condition in which the entire eye is ischemic?
Ocular ischemic syndrome (OIS)

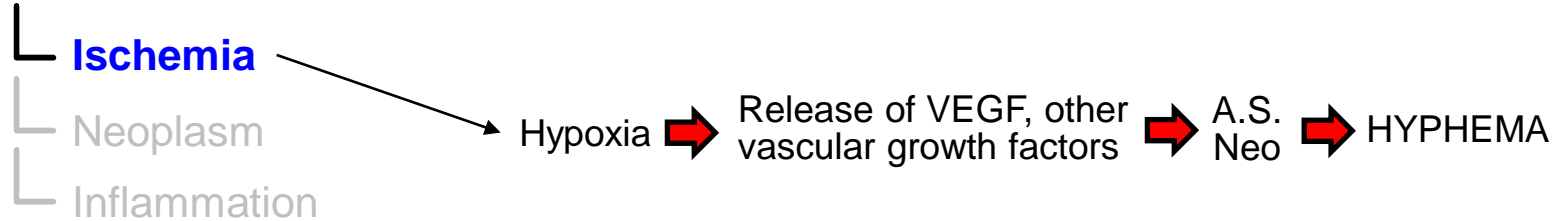
In OIS, where in the vascular tree is the occlusion found?

A

Spontaneous Hyphema

Adults

Children



Broadly speaking, ischemia of one of two 'structures' (note the 'scare quotes') is responsible for producing anterior segment neo. What are these structures?

The Retina, and the **Entire Eye**

What is the name of the condition in which the entire eye is ischemic?
Ocular ischemic syndrome (OIS)

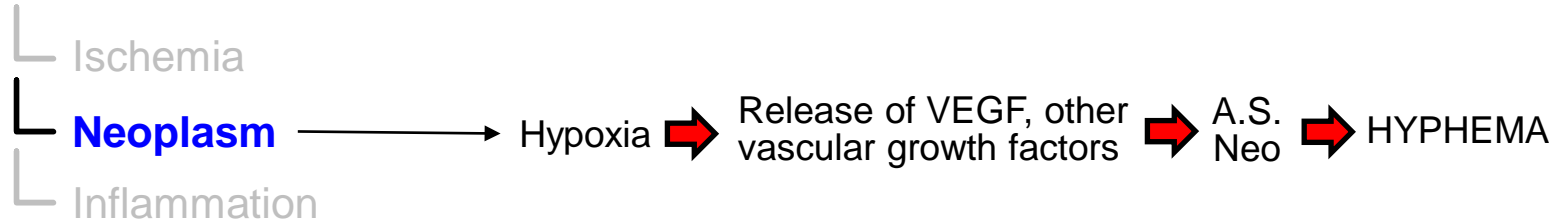
In OIS, where in the vascular tree is the occlusion found?
The ipsilateral carotid

Q

Spontaneous Hyphema

Adults

Children



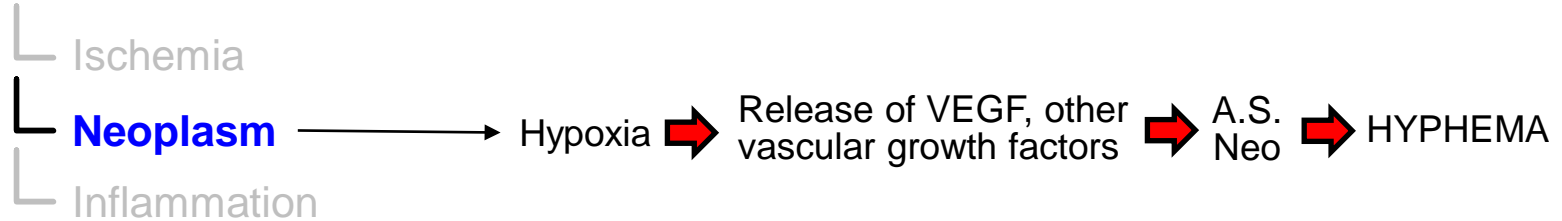
OK, I get how ischemia leads to hypoxia, but how does a neoplasm?

A

Spontaneous Hyphema

Adults

Children



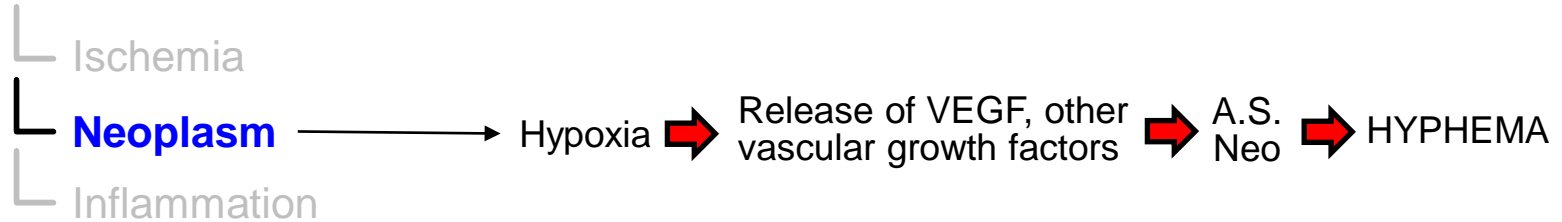
OK, I get how ischemia leads to hypoxia, but how does a neoplasm?
 Some neoplasms grow so rapidly that they end up outgrowing their blood supply. Tumor cells located at the fringe of the available blood supply may be hypoxic, which leads to their production and release of pro-vascular growth factors resulting in neo, and hyphema.

Q

Spontaneous Hyphema

Adults

Children

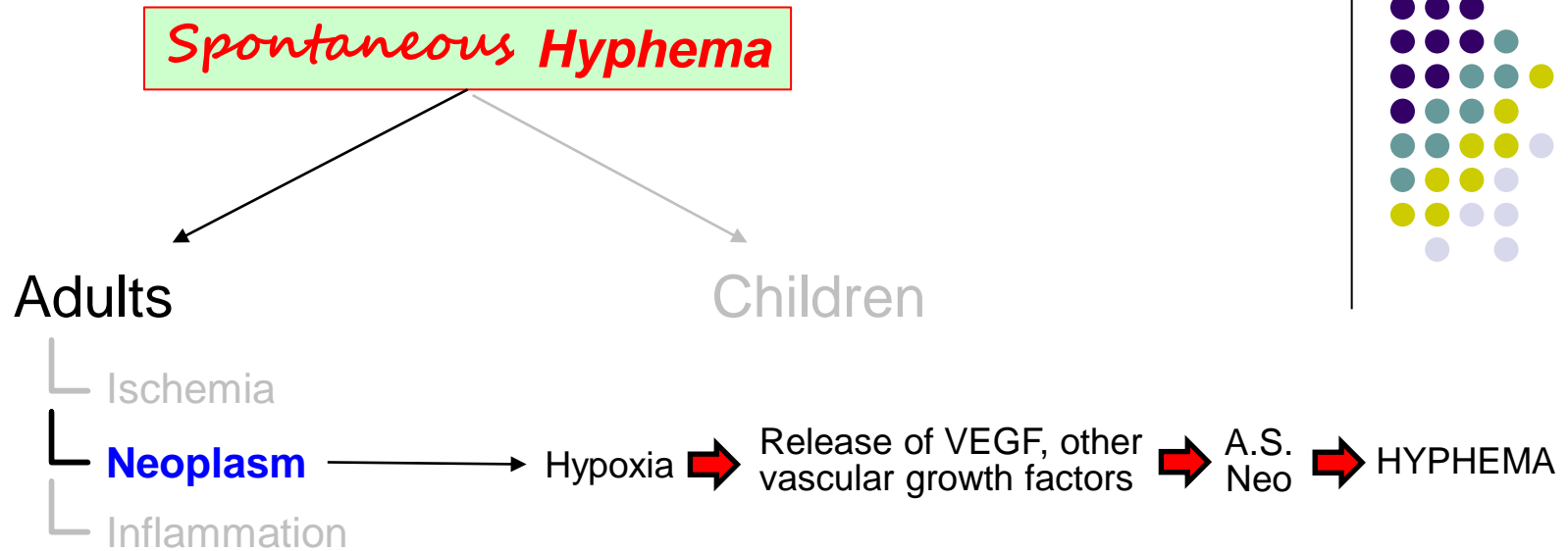


OK, I get how ischemia leads to hypoxia, but how does a neoplasm?

Some neoplasms grow so rapidly that they end up outgrowing their blood supply. Tumor cells located at the fringe of the available blood supply may be hypoxic, which leads to their production and release of pro-vascular growth factors resulting in neo, and hyphema.

Which neoplasm is perhaps best known to be associated with hyphema in adults?

A



OK, I get how ischemia leads to hypoxia, but how does a neoplasm?

Some neoplasms grow so rapidly that they end up outgrowing their blood supply. Tumor cells located at the fringe of the available blood supply may be hypoxic, which leads to their production and release of pro-vascular growth factors resulting in neo, and hyphema.

Which neoplasm is perhaps best known to be associated with hyphema in adults?

Uveal melanoma

Q

Spontaneous Hyphema

Adults

- └ Ischemia
- └ Neoplasm
- └ **Inflammation**

Children

Hypoxia → Release of VEGF, other vascular growth factors → A.S. Neo → HYPHEMA

How does inflammation lead to neo, and then hyphema?



A

Spontaneous Hyphema

Adults

Children

- └ Ischemia
- └ Neoplasm
- └ **Inflammation**

Hypoxia → Release of VEGF, other vascular growth factors → A.S. Neo → HYPHEMA

How does inflammation lead to neo, and then hyphema?

This has yet to be fully elucidated, and the *BCSC* books do not address the 'mechanism' question



Q

Spontaneous Hyphema

Adults

- └ Ischemia
- └ Neoplasm
- └ **Inflammation**

Children

Hypoxia → Release of VEGF, other vascular growth factors → A.S. Neo → HYPHEMA

How does inflammation lead to neo, and then hyphema?

This has yet to be fully elucidated, and the *BCSC* books do not address the 'mechanism' question

Which forms of uveitis are well-known to be associated with hyphema in adults?

--

--



A

Spontaneous Hyphema

Adults

Children

- └ Ischemia
- └ Neoplasm
- └ **Inflammation**

Hypoxia → Release of VEGF, other vascular growth factors → A.S. Neo → HYPHEMA

How does inflammation lead to neo, and then hyphema?

This has yet to be fully elucidated, and the *BCSC* books do not address the 'mechanism' question

Which forms of uveitis are well-known to be associated with hyphema in adults?

- Herpetic uveitides
- Fuch heterochromic iridocyclitis (FHI)



Q

Spontaneous Hyphema

Adults

- └ Ischemia
- └ Neoplasm
- └ **Inflammation**

Children

Hypoxia → Release of VEGF, other vascular growth factors → A.S. Neo → HYPHEMA

How does inflammation lead to neo, and then hyphema?

This has yet to be fully elucidated, and the *BCSC* books do not address the 'mechanism' question

Which forms of uveitis are well-known to be associated with hyphema in adults?

-- **Herpetic uveitides**

-- Fuchs heterochromic iridocyclitis (FHI)

Which herpetic uveitides are especially notorious for hyphema?

--

--



A

Spontaneous Hyphema

Adults

- └ Ischemia
- └ Neoplasm
- └ **Inflammation**

Children

Hypoxia → Release of VEGF, other vascular growth factors → A.S. Neo → HYPHEMA

How does inflammation lead to neo, and then hyphema?

This has yet to be fully elucidated, and the *BCSC* books do not address the 'mechanism' question

Which forms of uveitis are well-known to be associated with hyphema in adults?

--Herpetic uveitides

--Fuchs heterochromic iridocyclitis (FHI)

Which herpetic uveitides are especially notorious for hyphema?

--VZV

--HSV



Q

Spontaneous Hyphema

Adults

- └ Ischemia
- └ Neoplasm
- └ **Inflammation**

Children

Hypoxia → Release of VEGF, other vascular growth factors → A.S. Neo → HYPHEMA

How does inflammation lead to neo, and then hyphema?

This has yet to be fully elucidated, and the *BCSC* books do not address the 'mechanism' question

Which forms of uveitis are well-known to be associated with hyphema in adults?

--Herpetic uveitides

--Fuchs heterochromic iridocyclitis (FHI)

Which herpetic uveitides are especially associated with hyphema?

--VZV: Iris atrophy is...?

--HSV: Iris atrophy is...?

VZV and HSV both produce iris atrophy. In what key way do they differ in this regard?

A

Spontaneous Hyphema

Adults

- └ Ischemia
- └ Neoplasm
- └ **Inflammation**

Children

Hypoxia → Release of VEGF, other vascular growth factors → A.S. Neo → HYPHEMA

How does inflammation lead to neo, and then hyphema?

This has yet to be fully elucidated, and the *BCSC* books do not address the 'mechanism' question

Which forms of uveitis are well-known to be associated with hyphema in adults?

--**Herpetic uveitides**

--Fuchs heterochromic iridocyclitis (FHI)

Which herpetic uveitides are especially associated with hyphema?

--VZV: Iris atrophy is...**sectoral**

--HSV: Iris atrophy is...**diffuse**

VZV and HSV both produce iris atrophy. In what key way do they differ in this regard?

In VZV, the atrophy is **sectoral**, whereas in HSV it is **diffuse**

Q

Spontaneous Hypohemia

Who is the typical FHI pt?

--Herpetic uveitides

--**Fuch heterochromic iridocyclitis (FHI)**

EMA

A

Spontaneous Hypohemia

Who is the typical FHI pt?

A middle-aged adult

--Herpetic uveitides

--Fuch heterochromic iridocyclitis (FHI)

Q

Spontaneous Hypohemia

Who is the typical FHI pt?

A middle-aged adult

Is there a gender predilection?

--Herpetic uveitides

--**Fuch heterochromic iridocyclitis (FHI)**

A

Spontaneous Hypohemia

Who is the typical FHI pt?

A middle-aged adult

Is there a gender predilection?

No

--Herpetic uveitides

--**Fuch heterochromic iridocyclitis (FHI)**

Q

Spontaneous Hyphema

Who is the typical FHI pt?

A middle-aged adult

Is there a gender predilection?

No

What exam findings comprise the 'classic triad' of FHI? (Hyphema is not one of them.)

--Herpetic uveitides

--**Fuch heterochromic iridocyclitis (FHI)**

A

Spontaneous Hyphema

Who is the typical FHI pt?

A middle-aged adult

Is there a gender predilection?

No

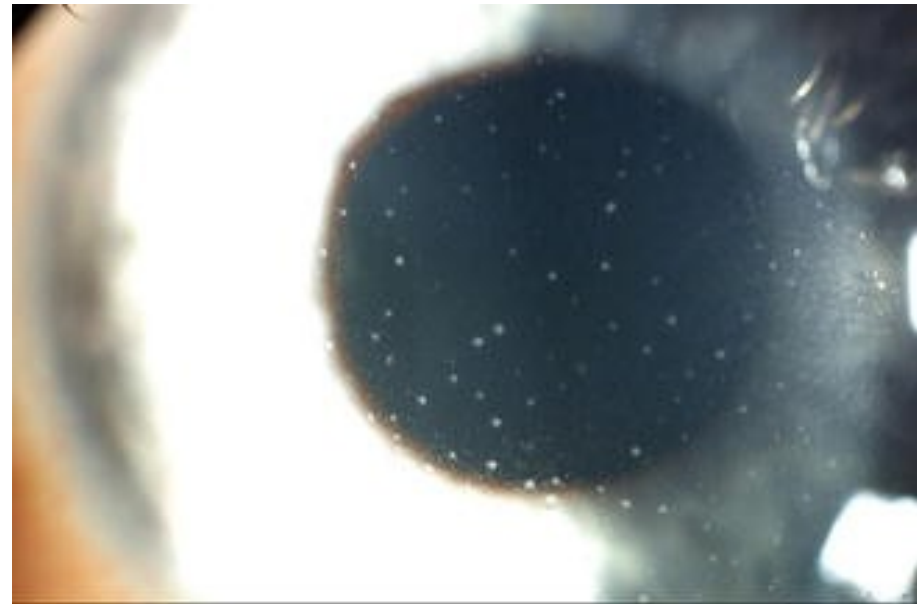
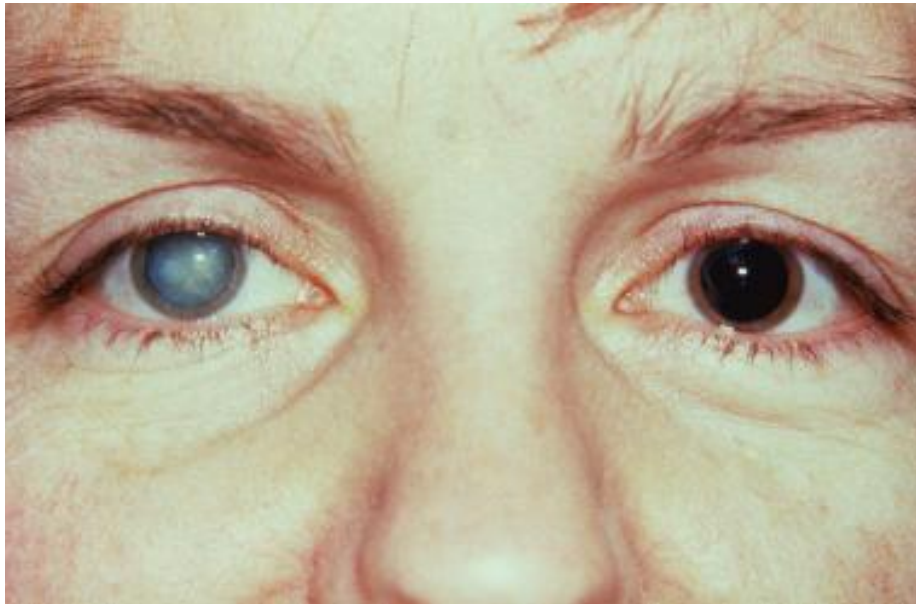
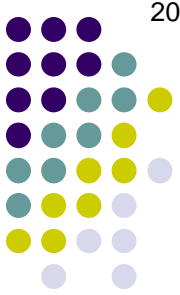
What exam findings comprise the 'classic triad' of FHI? (Hyphema is not one of them.)

Heterochromia iridis, cataract, and stellate KP

--Herpetic uveitides

--**Fuch heterochromic iridocyclitis (FHI)**

Spontaneous Hyphema



FHI: Heterochromia iridis, cataract, and stellate KP

Q

Spontaneous Hyphema

Who is the typical FHI pt?

A middle-aged adult

Is there a gender predilection?

No

What exam findings comprise the 'classic triad' of FHI? (Hyphema is not one of them.)

Heterochromia iridis, cataract, and stellate KP



Is the affected eye the **darker** eye, or the **lighter** eye?

--Herpetic uveitides

--Fuch heterochromic iridocyclitis (FHI)

A

Spontaneous Hyphema

Who is the typical FHI pt?

A middle-aged adult

Is there a gender predilection?

No

What exam findings comprise the 'classic triad' of FHI? (Hyphema is not one of them.)

Heterochromia iridis, cataract, and stellate KP



Is the affected eye the **darker** eye, or the **lighter** eye?
The lighter (most of the time; see U13 for details)

--Herpetic uveitides

--**Fuch heterochromic iridocyclitis (FHI)**

Spontaneous Hyphema



FHI

Q

Spontaneous Hyphema

Who is the typical FHI pt?

A middle-aged adult

Is there a gender predilection?

No

What exam findings comprise the 'classic triad' of FHI? (Hyphema is not one of them.)

Heterochromia iridis, cataract, and stellate KP

What other ophthalmic issue is common in FHI? (Again, hyphema ain't it.)

--Herpetic uveitides

--**Fuch heterochromic iridocyclitis (FHI)**

A

Spontaneous Hyphema

Who is the typical FHI pt?

A middle-aged adult

Is there a gender predilection?

No

What exam findings comprise the 'classic triad' of FHI? (Hyphema is not one of them.)

Heterochromia iridis, cataract, and stellate KP

What other ophthalmic issue is common in FHI? (Again, hyphema ain't it.)

Glaucoma—it develops in about 25-50% of cases

--Herpetic uveitides

--**Fuch heterochromic iridocyclitis (FHI)**

Q

Spontaneous Hyphema

Who is the typical FHI pt?

A middle-aged adult

Is there a gender predilection?

No

What exam findings comprise the 'classic triad' of FHI? (Hyphema is not one of them.)

Heterochromia iridis, cataract, and stellate KP

What other ophthalmic issue is common in FHI? (Again, hyphema ain't it.)

Glaucoma—it develops in about 25-50% of cases

What is the etiology of FHI?

--Herpetic uveitides

--**Fuch heterochromic iridocyclitis (FHI)**

Q/A

Spontaneous Hyphema

Who is the typical FHI pt?

A middle-aged adult

Is there a gender predilection?

No

What exam findings comprise the 'classic triad' of FHI? (Hyphema is not one of them.)

Heterochromia iridis, cataract, and stellate KP

What other ophthalmic issue is common in FHI? (Again, hyphema ain't it.)

Glaucoma—it develops in about 25-50% of cases

What is the etiology of FHI?

It is uncertain at this time. Four infectious entities have been suggested:

--?

--?

--?

--?

--Herpetic uveitides

Fuch heterochromic iridocyclitis (FHI)

A

Spontaneous Hyphema

Who is the typical FHI pt?

A middle-aged adult

Is there a gender predilection?

No

What exam findings comprise the 'classic triad' of FHI? (Hyphema is not one of them.)

Heterochromia iridis, cataract, and stellate KP

What other ophthalmic issue is common in FHI? (Again, hyphema ain't it.)

Glaucoma—it develops in about 25-50% of cases

What is the etiology of FHI?

It is uncertain at this time. Four infectious entities have been suggested:

--Toxoplasmosis

--HSV

--CMV

--Rubella

--Herpetic uveitides

--Fuch heterochromic iridocyclitis (FHI)

Q

Spontaneous Hyphema

Who is the typical FHI pt?

A middle-aged adult

Is there a gender predilection?

No

What exam findings comprise the 'classic triad' of FHI? (Hyphema is not one of them.)

Heterochromia iridis, cataract, and stellate KP

What other ophthalmic issue is common in FHI? (Again, hyphema ain't it.)

Glaucoma—it develops in about 25-50% of cases

What is the etiology of FHI?

It is uncertain at this time. Four infectious entities have been suggested:

--*Toxoplasmosis?*

--*HSV?*

--*CMV?*

--*Rubella?*

As of now, the preponderance of the evidence points to , but it remains unproven.

--Herpetic uveitides

Fuch heterochromic iridocyclitis (FHI)

A

Spontaneous Hyphema

Who is the typical FHI pt?

A middle-aged adult

Is there a gender predilection?

No

What exam findings comprise the 'classic triad' of FHI? (Hyphema is not one of them.)

Heterochromia iridis, cataract, and stellate KP

What other ophthalmic issue is common in FHI? (Again, hyphema ain't it.)

Glaucoma—it develops in about 25-50% of cases

What is the etiology of FHI?

It is uncertain at this time. Four infectious entities have been suggested:

--Toxoplasmosis

--HSV

--CMV

--**Rubella!**

As of now, the preponderance of the evidence points to rubella , but it remains unproven.

--Herpetic uveitides

--**Fuch heterochromic iridocyclitis (FHI)**

Q

Spontaneous Hyphema

Who is the typical FHI pt?

A middle-aged adult

Is there a gender predilection?

No

What exam findings comprise the 'classic triad' of FHI? (Hyphema is not one of them.)

Heterochromia iridis, cataract, and stellate KP

What other ophthalmic issue is common in FHI? (Again, hyphema ain't it.)

How well does FHI respond to steroid therapy?

--Herpetic uveitides

--**Fuch heterochromic iridocyclitis (FHI)**

A

Spontaneous Hyphema

Who is the typical FHI pt?

A middle-aged adult

Is there a gender predilection?

No

What exam findings comprise the 'classic triad' of FHI? (Hyphema is not one of them.)

Heterochromia iridis, cataract, and stellate KP

What other ophthalmic issue is common in FHI? (Again, hyphema ain't it.)

How well does FHI respond to steroid therapy?

Poorly--AC cell is notoriously difficult to eradicate in FHI (as is the anterior vitreous cell which frequently occurs)

--Herpetic uveitides

Fuch heterochromic iridocyclitis (FHI)

Q

Spontaneous Hyphema

Who is the typical FHI pt?

A middle-aged adult

Is there a gender predilection?

No

What exam findings comprise the 'classic triad' of FHI? (Hyphema is not one of them.)

Heterochromia iridis, cataract, and stellate KP

What other ophthalmic issue is common in FHI? (Again, hyphema ain't it.)

How well does FHI respond to steroid therapy?

Poorly--AC cell is notoriously difficult to eradicate in FHI (as is the anterior vitreous cell which frequently occurs)

If/when the inflammation fails to respond to topical steroids, should more aggressive therapies be pursued?

--Herpetic uveitides

Fuch heterochromic iridocyclitis (FHI)

A

Spontaneous Hyphema

Who is the typical FHI pt?

A middle-aged adult

Is there a gender predilection?

No

What exam findings comprise the 'classic triad' of FHI? (Hyphema is not one of them.)

Heterochromia iridis, cataract, and stellate KP

What other ophthalmic issue is common in FHI? (Again, hyphema ain't it.)

How well does FHI respond to steroid therapy?

Poorly--AC cell is notoriously difficult to eradicate in FHI (as is the anterior vitreous cell which frequently occurs)

If/when the inflammation fails to respond to topical steroids, should more aggressive therapies be pursued?

Generally no. In fact, most pts require no anti-inflammatory tx of any sort (including steroids). Instead, the pt should be monitored for the development of glaucoma and cataract.

--Herpetic uveitides

Fuch heterochromic iridocyclitis (FHI)

Q

Spontaneous Hyphema

Adults

- └ Ischemia
- └ Neoplasm
- └ Inflammation

Children

- └ ?
- └ ?
- └ ?

What three conditions are most commonly associated with spontaneous hyphema in children?



A

Spontaneous Hyphema

Adults

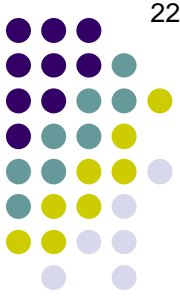
- └ Ischemia
- └ Neoplasm
- └ Inflammation

Children

- └ JXG
- └ Rb
- └ Leukemia

What three conditions are most commonly associated with spontaneous hyphema in children?





Q

Spontaneous Hyphema

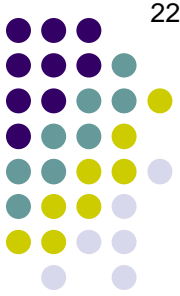
Adults

- └ Ischemia
- └ Neoplasm
- └ Inflammation

Children

- └ JXG

What does JXG stand for?



A

Spontaneous Hyphema

Adults

- └ Ischemia
- └ Neoplasm
- └ Inflammation

Children

└ **JXG**

What does JXG stand for?
Juvenile xanthogranuloma



Q

Spontaneous Hyphema

Adults

- └ Ischemia
- └ Neoplasm
- └ Inflammation

Children

└ JXG

What does JXG stand for?
Juvenile xanthogranuloma

In three words, what sort of condition is it?
It is a



A

Spontaneous Hyphema

Adults

- └ Ischemia
- └ Neoplasm
- └ Inflammation

Children

└ JXG

What does JXG stand for?
Juvenile xanthogranuloma

In three words, what sort of condition is it?
It is a **nonneoplastic histiocytic proliferation**



Q

Spontaneous Hyphema

Adults

- └ Ischemia
- └ Neoplasm
- └ Inflammation

Children

└ JXG

What does JXG stand for?
Juvenile xanthogranuloma

In three words, what sort of condition is it?
It is a **nonneoplastic histiocytic proliferation**

At what age does JXG present?



A

Spontaneous Hyphema

Adults

- └ Ischemia
- └ Neoplasm
- └ Inflammation

Children

└ **JXG**

What does JXG stand for?
Juvenile xanthogranuloma

In three words, what sort of condition is it?
It is a **nonneoplastic histiocytic proliferation**

At what age does JXG present?
The majority before age 1 year, and almost all by age 2



Q

Spontaneous Hyphema

Adults

- └ Ischemia
- └ Neoplasm
- └ Inflammation

Children

└ JXG

What does JXG stand for?
Juvenile xanthogranuloma

In three words, what sort of condition is it?
It is a **nonneoplastic histiocytic proliferation**

At what age does JXG present?
The majority before age 1 year, and almost all by age 2

How does JXG usually present? (Hint: It's not ophthalmic)



A

Spontaneous Hyphema

Adults

- └ Ischemia
- └ Neoplasm
- └ Inflammation

Children

└ JXG

What does JXG stand for?
Juvenile xanthogranuloma

In three words, what sort of condition is it?
It is a **nonneoplastic histiocytic proliferation**

At what age does JXG present?
The majority before age 1 year, and almost all by age 2

How does JXG usually present? (Hint: It's not ophthalmic)
As skin papules



JXG: Skin papules. The orangish color is classic

Q

Spontaneous Hyphema

Adults

- └ Ischemia
- └ Neoplasm
- └ Inflammation

Children

└ JXG

What does JXG stand for?
Juvenile xanthogranuloma

In three words, what sort of condition is it?
It is a **nonneoplastic histiocytic proliferation**

At what age does JXG present?
The majority before age 1 year, and almost all by age 2

How does JXG usually present? (Hint: It's not ophthalmic)
As skin papules

*When **iris** JXG nodules are present, is it uni-, or bilaterally?*



A

Spontaneous Hyphema

Adults

- └ Ischemia
- └ Neoplasm
- └ Inflammation

Children

└ JXG

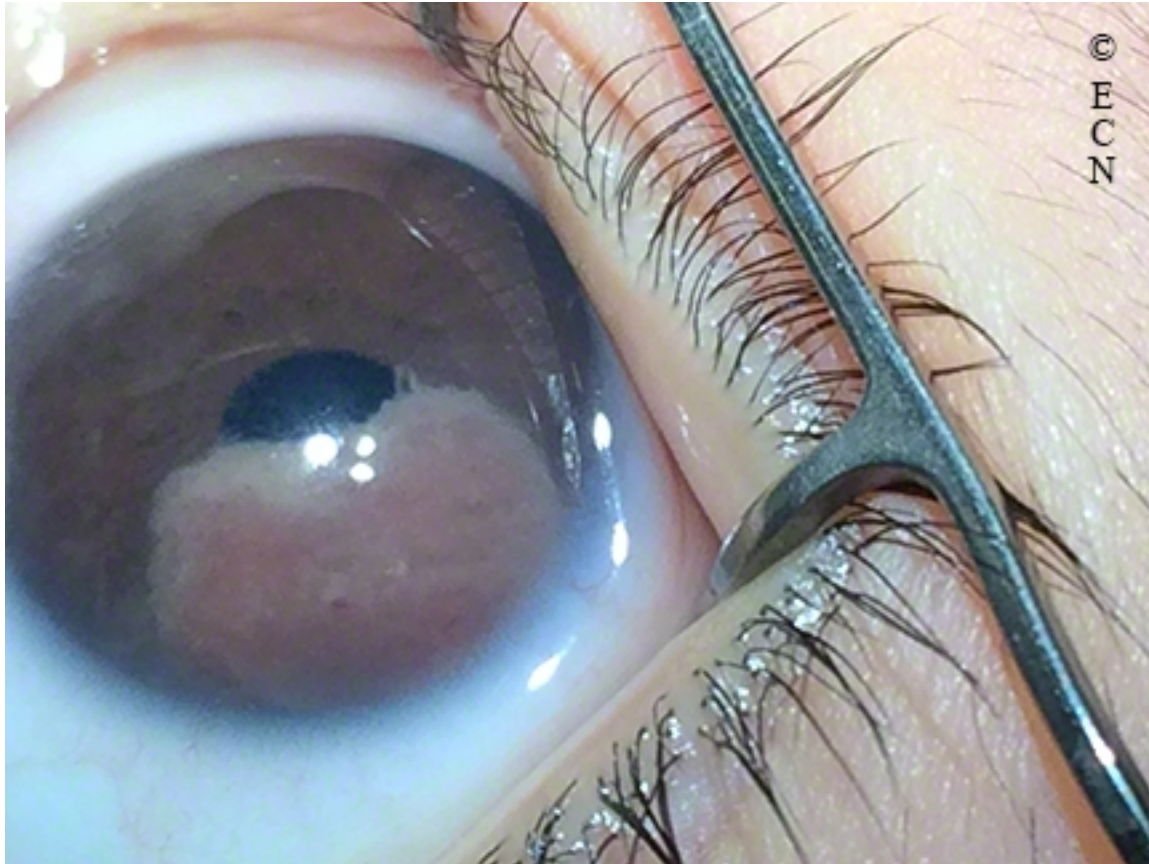
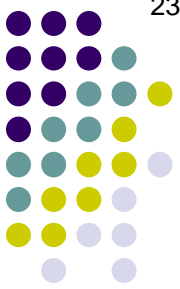
What does JXG stand for?
Juvenile xanthogranuloma

In three words, what sort of condition is it?
It is a **nonneoplastic histiocytic proliferation**

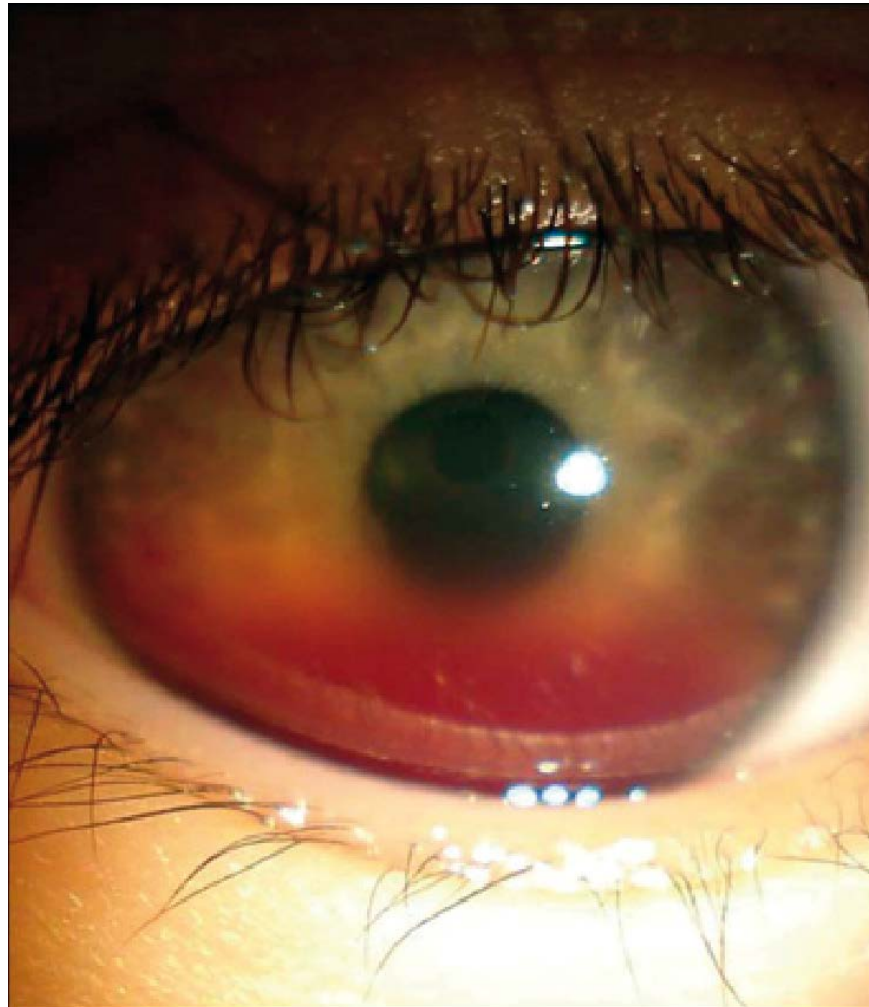
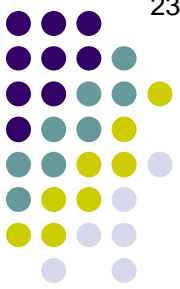
At what age does JXG present?
The majority before age 1 year, and almost all by age 2

How does JXG usually present? (Hint: It's not ophthalmic)
As skin papules

*When **iris** JXG nodules are present, is it uni-, or bilaterally?*
Unilaterally



JXG: Iris lesion



JXG: Spontaneous hyphema

Q

Spontaneous Hyphema

Adults

- └ Ischemia
- └ Neoplasm
- └ Inflammation

Children

- └ JXG

What does JXG stand for?

In addition to spontaneous hyphema, in what three ways are JXG iris nodules clinically significant?

--
--
--

When iris JXG nodules are present, is it uni-, or bilaterally?
Unilaterally



A

Spontaneous Hyphema

Adults

- └ Ischemia
- └ Neoplasm
- └ Inflammation

Children

└ JXG

What does JXG stand for?

In addition to spontaneous hyphema, in what three ways are JXG iris nodules clinically significant?

- They are in the DDx as a 'masquerade syndrome' in peds uveitis
- If enough are present, they will cause heterochromia iridis
- They can result in severe glaucoma

When iris JXG nodules are present, is it uni-, or bilaterally?

Unilaterally



Q

Spontaneous Hyphema

Adults

- └ Ischemia
- └ Neoplasm
- └ Inflammation

Children

└ JXG

What does JXG stand for?

In addition to spontaneous hyphema, in what three ways are JXG iris nodules clinically significant?

- They are in the DDx as a 'masquerade syndrome' in peds uveitis
- If enough are present, they will cause heterochromia iridis
- They can result in severe glaucoma

Should they be removed surgically?

When iris JXG nodules are present, is it uni-, or bilaterally?

Unilaterally



A

Spontaneous Hyphema

Adults

- └ Ischemia
- └ Neoplasm
- └ Inflammation

Children

└ JXG

What does JXG stand for?

In addition to spontaneous hyphema, in what three ways are JXG iris nodules clinically significant?

- They are in the DDx as a 'masquerade syndrome' in peds uveitis
- If enough are present, they will cause heterochromia iridis
- They can result in severe glaucoma

Should they be removed surgically?

Only if the glaucoma is uncontrollable

When iris JXG nodules are present, is it uni-, or bilaterally?

Unilaterally





Q

Spontaneous Hyphema

Adults

- └ Ischemia
- └ Neoplasm
- └ Inflammation

Children

- └ JXG

What does JXG stand for?

In addition to spontaneous hyphema, in what three ways are JXG iris nodules clinically significant?

- They are in the DDx as a 'masquerade syndrome' in peds uveitis
- If enough are present, they will cause heterochromia iridis
- They can result in severe glaucoma

Should they be removed surgically?

Only if the glaucoma is uncontrollable

What is the natural history of the disease?

When iris JXG nodules are present, is it uni-, or bilaterally?

Unilaterally

Q/A

Spontaneous Hyphema

Adults

- └ Ischemia
- └ Neoplasm
- └ Inflammation

Children

- └ JXG

What does JXG stand for?

In addition to spontaneous hyphema, in what three ways are JXG iris nodules clinically significant?

- They are in the DDx as a 'masquerade syndrome' in peds uveitis
- If enough are present, they will cause heterochromia iridis
- They can result in severe glaucoma

Should they be removed surgically?

Only if the glaucoma is uncontrollable

What is the natural history of the disease?

JXG is self-limited, usually resolving by age 2

When iris JXG nodules are present, is it uni-, or bilaterally?

Unilaterally



A

Spontaneous Hyphema

Adults

- └ Ischemia
- └ Neoplasm
- └ Inflammation

Children

- └ JXG

What does JXG stand for?

In addition to spontaneous hyphema, in what three ways are JXG iris nodules clinically significant?

- They are in the DDx as a 'masquerade syndrome' in peds uveitis
- If enough are present, they will cause heterochromia iridis
- They can result in severe glaucoma

Should they be removed surgically?

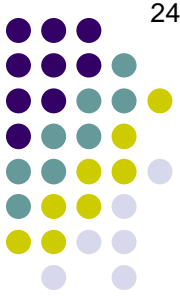
Only if the glaucoma is uncontrollable

What is the natural history of the disease?

JXG is self-limited, usually resolving by age 5 years

When iris JXG nodules are present, is it uni-, or bilaterally?

Unilaterally



Q

Spontaneous Hyphema

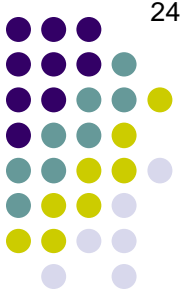
Adults

- └ Ischemia
- └ Neoplasm
- └ Inflammation

Children

- └ JXG
- └ **Rb**
- └ Leukemia

What does Rb stand for in this context?



A

Spontaneous Hyphema

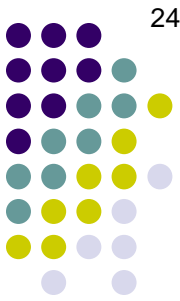
Adults

- └ Ischemia
- └ Neoplasm
- └ Inflammation

Children

- └ JXG
- └ **Rb**
- └ Leukemia

What does Rb stand for in this context?
Retinoblastoma



Q

Spontaneous Hyphema

Adults

- └ Ischemia
- └ Neoplasm
- └ Inflammation

Children

- └ JXG
- └ **Rb**
- └ Leukemia

What does Rb stand for in this context?
Retinoblastoma

The BCSC doesn't have too much to say about hyphema in Rb, save the following two points:
--It is [common vs uncommon]



A

Spontaneous Hyphema

Adults

- └ Ischemia
- └ Neoplasm
- └ Inflammation

Children

- └ JXG
- └ **Rb**
- └ Leukemia

What does Rb stand for in this context?
Retinoblastoma

The BCSC doesn't have too much to say about hyphema in Rb, save the following two points:
--It is uncommon

Q

Spontaneous Hyphema

Adults

- └ Ischemia
- └ Neoplasm
- └ Inflammation

Children

- └ JXG
- └ **Rb**
- └ Leukemia

What does Rb stand for in this context?
Retinoblastoma

The BCSC doesn't have too much to say about hyphema in Rb, save the following two points:

- It is uncommon
- It is more likely to occur in children than 5 years of age

[> or <]





A

Spontaneous Hyphema

Adults

- └ Ischemia
- └ Neoplasm
- └ Inflammation

Children

- └ JXG
- └ **Rb**
- └ Leukemia

What does Rb stand for in this context?
Retinoblastoma

The BCSC doesn't have too much to say about hyphema in Rb, save the following two points:

- It is uncommon
- It is more likely to occur in children < than 5 years of age



Spontaneous Hyphema

Adults

- └ Ischemia
- └ Neoplasm
- └ Inflammation

Children

- └ JXG
- └ **Rb**
- └ Leukemia

What does Rb stand for in this context?
Retinoblastoma

The BCSC doesn't have too much to say about hyphema in Rb, save the following two points:

- It is uncommon
- It is more likely to occur in children < than 5 years of age

For more on Rb, see slide-set R2

Q

Spontaneous Hyphema

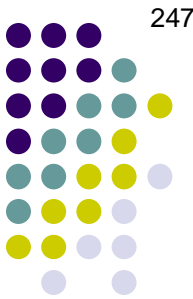
Adults

- └ Ischemia
- └ Neoplasm
- └ Inflammation

Children

- └ JXG
- └ Rb
- └ **Leukemia**

Where does leukemia rank among childhood malignancies in terms of incidence?



A

Spontaneous Hyphema

Adults

- └ Ischemia
- └ Neoplasm
- └ Inflammation

Children

- └ JXG
- └ Rb
- └ **Leukemia**

Where does leukemia rank among childhood malignancies in terms of incidence?

It is #1 by a mile



Q

Spontaneous Hyphema

Adults

- └ Ischemia
- └ Neoplasm
- └ Inflammation

Children

- └ JXG
- └ Rb
- └ **Leukemia**

Where does leukemia rank among childhood malignancies in terms of incidence?

It is #1 by a mile

Re its histology: Which form of leukemia is most common in children?



A

Spontaneous Hyphema

Adults

- └ Ischemia
- └ Neoplasm
- └ Inflammation

Children

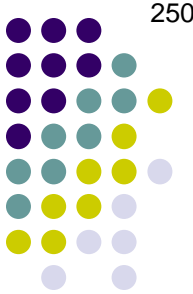
- └ JXG
- └ Rb
- └ **Leukemia**

Where does leukemia rank among childhood malignancies in terms of incidence?

It is #1 by a mile

Re its histology: Which form of leukemia is most common in children?

Acute lymphocytic (ALL)



Q

Spontaneous Hyphema

Adults

- └ Ischemia
- └ Neoplasm
- └ Inflammation

Children

- └ JXG
- └ Rb
- └ **Leukemia**



Where does leukemia rank among childhood malignancies in terms of incidence?

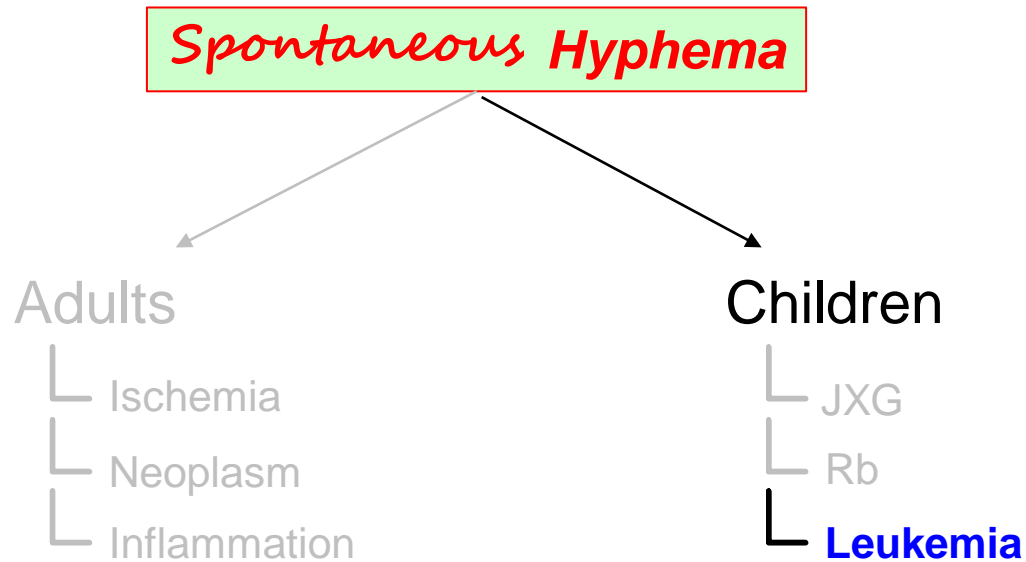
It is #1 by a mile

Re its histology: Which form of leukemia is most common in children?

Acute lymphocytic (ALL)

What other anterior-chamber condition is a well-known manifestation of leukemia?

A



Where does leukemia rank among childhood malignancies in terms of incidence?

It is #1 by a mile

Re its histology: Which form of leukemia is most common in children?

Acute lymphocytic (ALL)

What other anterior-chamber condition is a well-known manifestation of leukemia?

Pseudohypopyon

Spontaneous Hyphema

Adults

- └ Ischemia
- └ Neoplasm
- └ Inflammation

Children

- └ JXG
- └ Rb
- └ **Leukemia**

Where does leukemia rank among childhood malignancies in terms of incidence?

It is #1 by a mile

Re its histology: Which form of leukemia is most common in children?

Acute lymphocytic (ALL)

What other anterior-chamber condition is a well-known manifestation of leukemia?

Pseudohypopyon

For more on ophthalmic manifestations of pediatric leukemia, see slide-set P20

