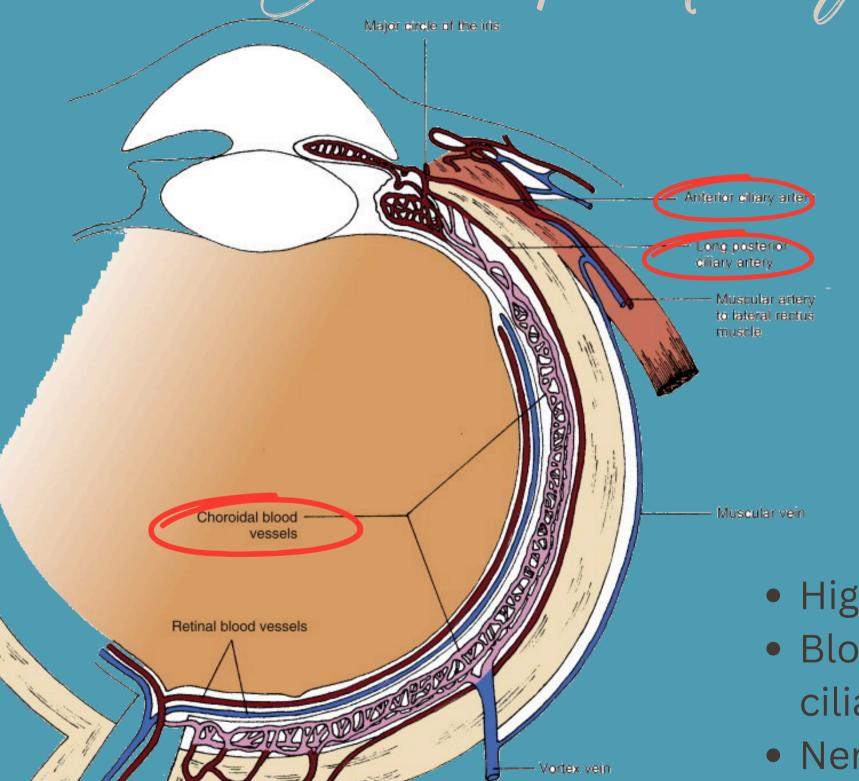


Kefla G. Brown, OD, MS

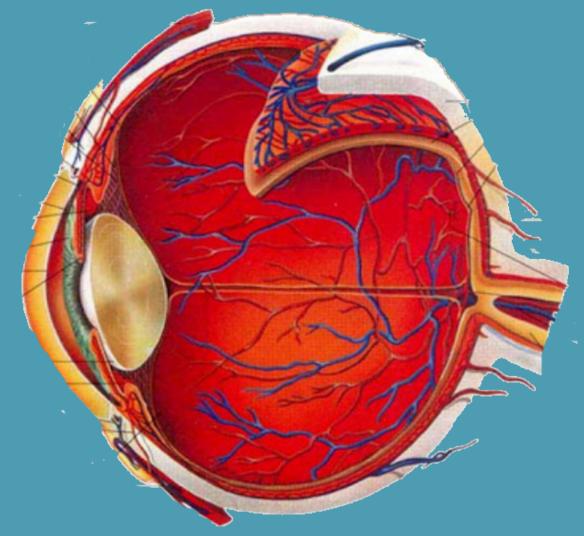


Basic Anatomy: Uveal Tract



ong posterior ciliary artery

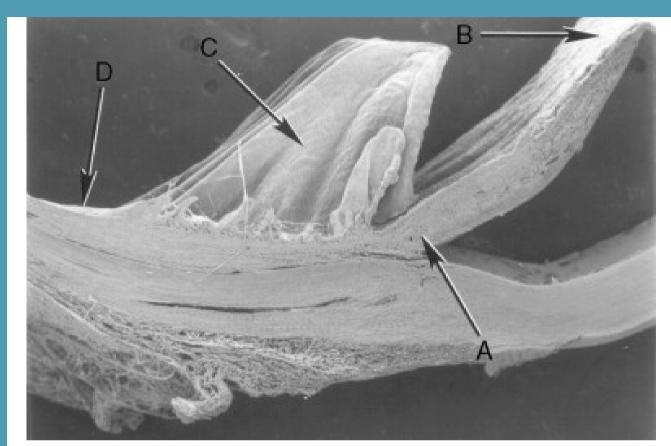
Central retinal artery.





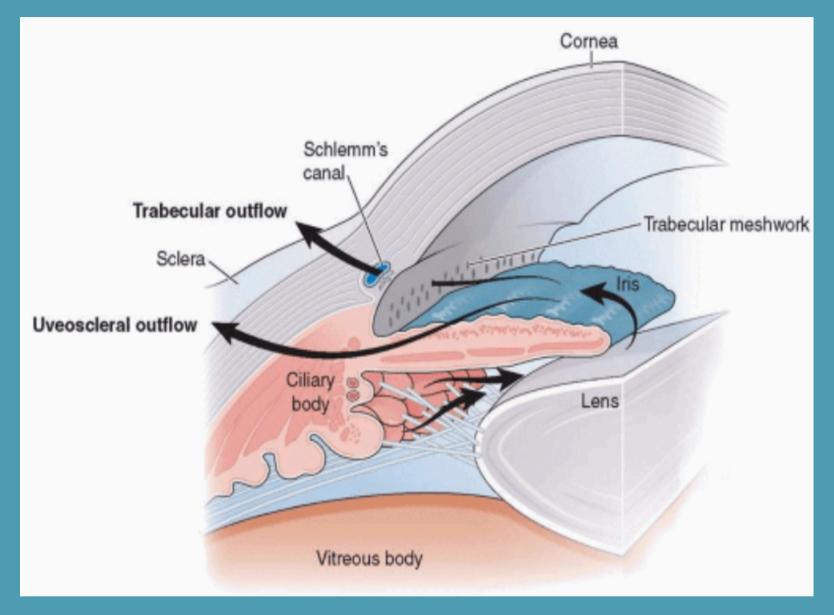
- Blood supply from the anterior and posterior ciliary arteries
- Nerve supply from anterior and posterior ciliary nerves
- High concentration of dendritic melanocytes

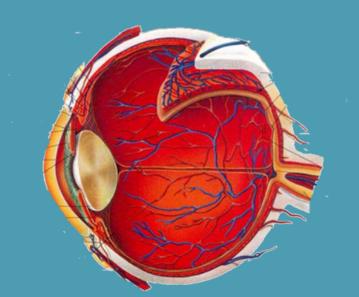
Ensic Anatomy: Meal Tract



The anterior uvea consists of the iris and ciliary body. Base of the iris (A), iridal pupillary margins (B), and ciliary body: pars plicata ciliaris (C) and pars plana ciliaris (D). SEM, 25×.

(Courtesy of Dr Don Samuelson, University of Florida.)





- Anterior Uveal Tract: Iris & Ciliary Body
- Posterior Uveal Tract: Choroid

Classification / Momenclature

LOCATION

Anterior

Intermediate

Posterior

Granulomatous

KERATIC PECIDITATE

Non-Granulomatous

Unilateral

LATERALITY

Bilateral

Acute

CHRONICITY

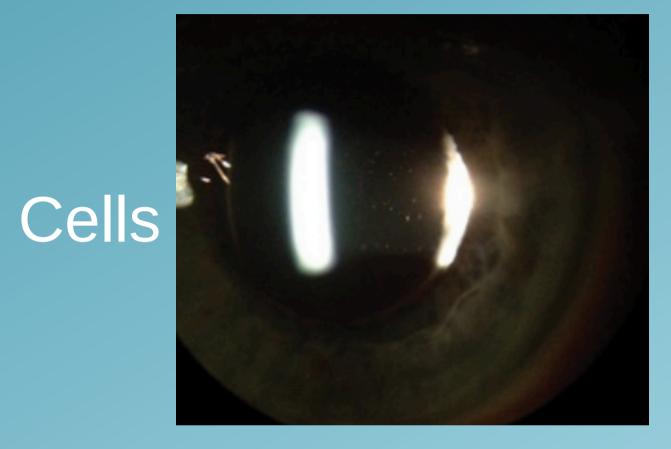
Chronic

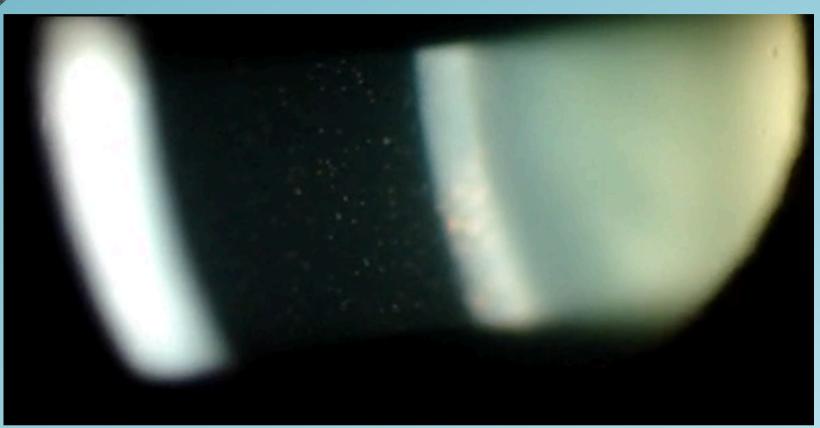
Painless

SYMPTOMS

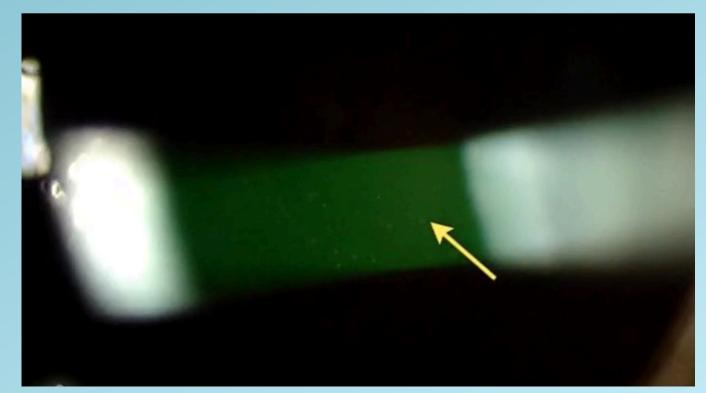
Painful

Signs





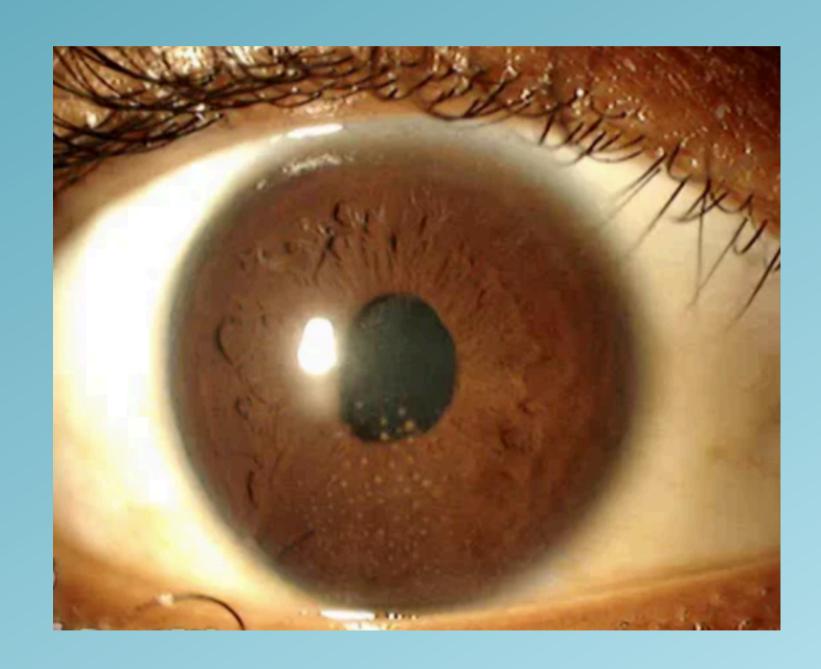
Flare

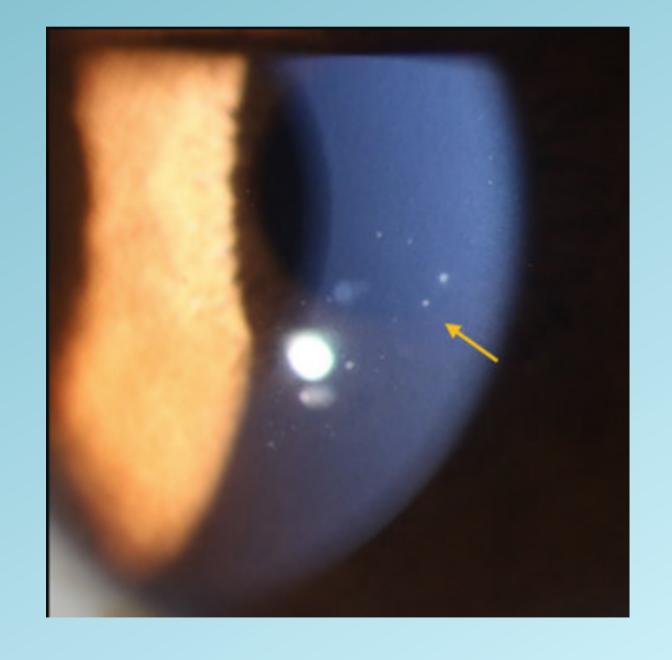






Signal Si





Granulomatous

Non - Granulomatous

Symptoms

Pain
Photophobia
Blurry Vision
Conjunctival injection (limbal flush)





Dinghosis

Bilateral Uveitis lab work up

Only done if specific condition suspected

ID	Name
LAB547	Sedimentation rate, manual
LAB2610	HLA B27 ANTIGEN
LAB179	ANGIOTENSIN CONVERTING ENZYME
LAB147	ANA PROFILE I (SCREEN)
LAB494	RPR *
LAB866	FTA ANTIBODIES, IGG AND IGM 💥
NUR559	POCT TB Skin Test 🕌
IMG34	X-Ray Chest 1 View 🕌
LAB206	Rheumatoid factor
LAB62	CK
LAB149	C-REACTIVE PROTEIN
LAB1748	CBC Auto Differential
LAB17	Comprehensive Metabolic Panel



Etology

"The majority of childhood uveitis syndromes are secondary to localized or generalized disorders of control of inflammation"*

- Idiopathic
- Traumatic
- Infectious
 - Histoplasmosis
 - Toxoplasmosis

- ANA +
- AutoimmuneDisease
 - Ulcerative Colitis/ Crohn's
 - o JIA





Trentment

- Topical Steroids
- Oral Steroids
- Systemic
 - Methotrexate
 - Biologics

- Co-management
- Side effects
- Long termconsiderations



MSOS

Case #1

4y.o. African-American Male

CC: Excessive blinking; "Just diagnosed with arthritis"

VA: 20/30; 20/30

lit Lamp and Fundus Exam					
External Exam					
	Right	Left			
External	Normal	Normal			
Slit Lamp Exam					
•	Right	Left			
Lids/Lashes	Normal	Normal			
Conjunctiva/Sclera	White and quiet	White and quiet			
Comea	Clear	Clear			
Anterior Chamber	2-3+ Cell	1+ Cell			
Iris	Round and reactive	Round and reactive			
Lens	Clear	Clear			

Diagnosis: Bilateral anterior uveitis (presumed etiology: JIA)

Treatment: Prednisolone Acetate 1% (PA), OU, Q.I.D.

Visit#2 (1 week):

- Decreased inflamation (1+ cell OD/ quiet OS)
- PA decreased to T.I.D. OU

Visit #3 (1 week):

- Decreased inflamation (Trace cell OD/ quiet OS)
- PA continued T.I.D. OU

Visit #4 (2 weeks):

- Stable
- Continued PA T.I.D OU (RTC 1 month)

Visit #5 (2 months)

Hospitalized for 5 weeks --> dx with Chronic Inflammatory Demyelinating Polyneuropathy (CIPD)

Next 6 months:

- Extended taper of PA
- Uveitis completely quiet

One year later: Updated diagnosis



Case #2

17y.o. Caucasian Male

- CC: Hit in the eye by phone charging cord that morning
- VA sc: OD 20/125 OS 20/200; PH : OD 20/40 OS20/20
- Pupils: OD 6-6 (-)DC No reaction (-) APD
 OS 4-3 (+)DC 4+/4+ (-) APD
- EOMs FROM OU

Slit Lamp Exam

	Right	Left
Lids/Lashes	erythema and mild edema at	Normal
	lateral aspect of upper lid	
Conjunctiva/Sclera	2+ Injection	White and quiet
Comea	Clear	Clear
Anterior Chamber	hyphema, 2+ Flare, 3+ Cell	Deep and quiet
Iris	hyphema	Round and reactive
Lens	Clear	Clear

Fundus Exam

	Right	Left
Vitreous	Normal	Normal
Disc	Normal, No pallor, no edema, no coloboma, no hypoplasia	Normal, No pallor, no edema, no coloboma, no hypoplasia
C/D Ratio	0.2	0.2
Macula	Normal, (+) foveal reflex	Normal, (+) foveal reflex
Vessels	Normal, Normal course and caliber, No tortuosity	Normal, Normal course and caliber, No tortuosity
Periphery	Normal, Flat & intact 360°, no holes, tears, RD	Normal, Flat & intact 360°, no holes, tears, RD



Assessment & Plan

1. Uveitis after traumatic injury of eye

- prednisoLONE acetate 1 %: 1 gtt, OD Q.I.D.
- homatropine 5 %: 1 gtt OD B.I.D.

2. Traumatic hyphema of left eye, initial encounter

- -Strict bedrest with bathroom privileges only; head elevated at 45 degrees
- -No NSAIDS
- 3. Retinal Health intact OU
- Return to clinic immediately with any new spontaneous flashes of light, a vail of gray, black or other color come over vision, or any new floaters

Visit #2: 1 Day f/u:

- Case #2, cont.
- VA sc: 20/125 OU PH : OD 20/25 + OU
- Pupils: OD 5-5 Minimal Reaction (-) APD OS 5-3 (+)DC 4+/4+ (-) APD
- IOP: 14 mmHg OD



Visit #2 - Assessment /Plan

- 1. Uveitis after traumatic injury of eye --> same cell, no flare
- -Increase prednisolone acetate 1% drops to every 2 hours while awake (OD) today, then back to every 4 hours tomorrow
- -Start cyclopentolate 1% twice daily, right eye, when available (OD)
- 2. Traumatic hyphema of left eye, subsequent encounter --> resolving
- -Decrease in blood in A/C (no 8 ball hyphema, no layering; no re-bleed)
- -Continue strict bedrest with bathroom privileges; head at 45 degree incline; avoid valsalva maneuvers

RTC in 1 day for retina, IOP, uveitis check, sooner as needed

Visit #3: 1 Day f/u:

Case #2, cont.

• VA cc: 20/20 OD

Pupils: "OD pupil is sluggish, dilated but reactive to light."

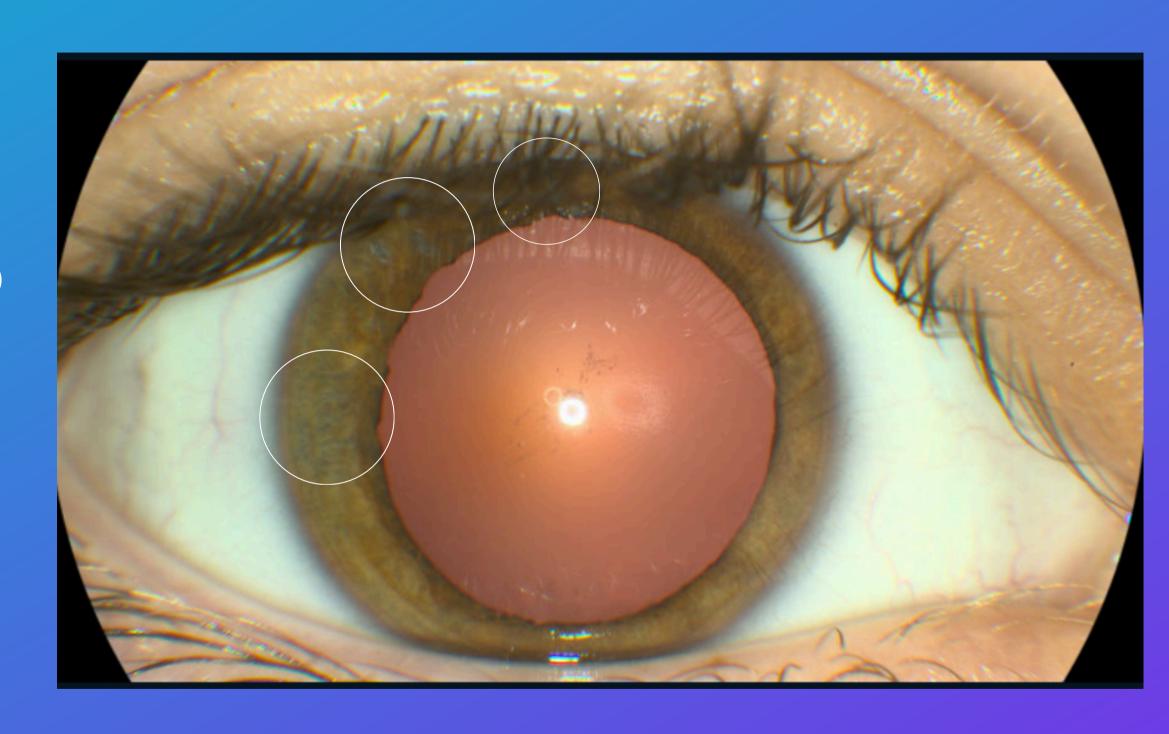
- IOP: 15 mmHg OD
- SLE: "min blod, Trace Cell, no flare"
- Bedrest maintained

Visit#4: 6 Day f/u:

- VA cc: 20/25 OD 20/20 OS
- Pupils: OD 5-4.5 Sluggish Reaction (-) APD OS 4-3 (+)DC 4+/4+ (-) APD
- IOP: 8 mmHg OD
- SLE: 1+ Cell; NO HYPHEMA; no flare
- A/P 1. PA increased to 4 times daily.
- ok to resume normal activities with care and caution
- RTC 1 week

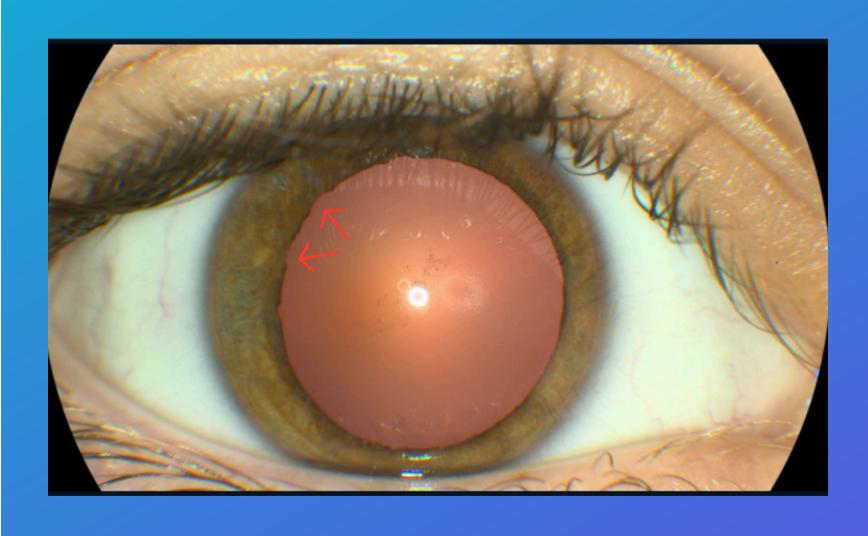
Visit #5: 1 Week f/u:

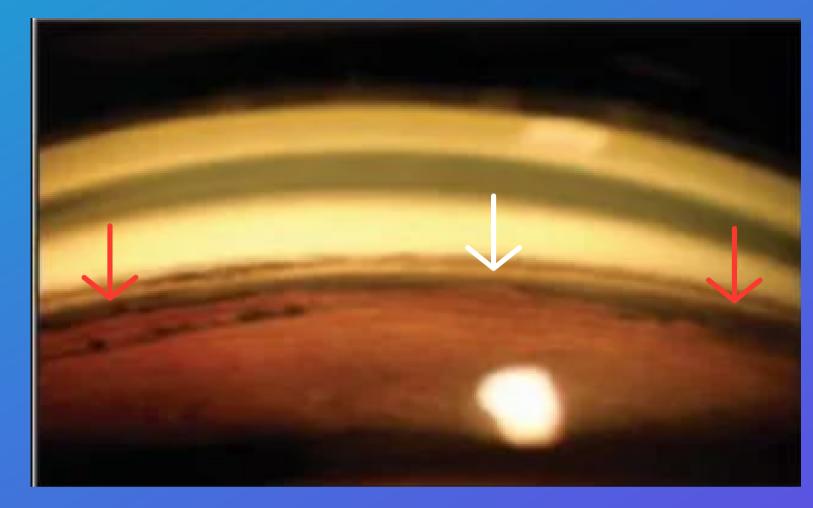
- BCVA: 20/20 OD (-1.25DS)
- Pupil:
 - \circ OD 6-5.5 (+) DC 2+/2+ (-) APD
 - \circ OS 4-3 (+) DC 4+/4+ (-) APD
- IOP OD: 12 mmHg
- 1+ Cell; no hyphema
- TRANSILLUMINATION DEFECTS



Referral to Glaucoma OMD:

- Angle Recession 360 degrees
- SphincterTear
- Transillumination Defects





A/P:

- Traumatic mydriasis OD with Iris sphincter tears and transillumination defects of the right iris
- Iris unable to constrict fully with light
- Increased risk of photosensitivity
- No acute surgical intervention today
- Could be a candidate for medical therapy if photosensitivity becomes bothersome

RTC 3 months with HVF24-2, OCT-RNFL, gonio, pachymetry

Case #3

6 y.o. Caucasian Female

- CC: Failed vision screening at PCP
- VA sc: OD 20/20 OS 20/40
- Pupils: Photopic:
 - OD 7 4 (+)DC 4+/4+ (-) APD
 - OS 5 4 (+)DC 1-/1- (-) APD
- Pupils: Scotopic:
 - \circ OD 8 6 (+)DC 4+/4+ (-) APD
 - OS 5 4 (+)DC 1-/1- (-) APD
- EOMs FROM OU
- Ishihara Normal, OU

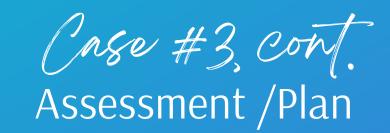
- Stereopsis: Global 125" Contour 50"
- Worth 4 Dot: Distance Suppression OS; near Fusion
- Bruckners: (+) anisocoria (right>left); dim reflex, left eye
- Cover Test: Ortho at distance; 10 XP at near (NPC: TN)

Cycloplegic refraction:

OD +0.50 sph

OS +2.50 sph

Slit Lamp and Fundus Exam				
External Exam				
	Right	Left		
External	Normal	Normal		
Palpebral fissure	11 mm	10 mm		
MRD1	3 mm	1 mm		
MRD2	5 mm	5 mm		
Superior scleral show	0 mm	0 mm		
Inferior scleral show	0 mm	0 mm		
Slit Lamp Exam				
	Right	Left		
Lids/Lashes	Normal	Normal		
Conjunctiva/Sclera	White and quiet	White and quiet		
Comea	Clear	Clear		
Anterior Chamber	Deep and quiet	2+ Flare		
Iris	Round and reactive	Posterior synechiae, Posterior synechiae 360 degrees		
		1: Posterior synechiae 360 degrees		
Lens	Clear	Anterior cortical changes/ diffuse mild opacity		
Fundus Exam				
	Right	Left		
Vitreous	Normal	Normal		
Disc	Normal, No pallor, no edema, no coloboma, no hypoplasia	Grossly Intact		
Macula	Normal, (+) foveal reflex	Grossly intact		
Vessels	Normal, Normal course and caliber, No tortuosity	Grossly intact		
Periphery	Normal, Flat & intact 360°, no holes, tears, RD	Grossly Intact		
Limited view, OS secondary to small pupil and lens opacity				



1. Chronic uveitis of left eye as evidenced by flare in anterior chamber and extensive posterior synechiae

Lab work up:

Sedimentation rate; HLA B27 Antigen; Angiotensin Converting Enzyme; ANA; Rheumatoid Factor; CK;

CRP; CBC auto differential; CMP

- Start treating with prednisolone acetate 1 %: 1 gtt, OS q. 4 hours

2. Posterior synechiae (iris), left eye

- -(+) significant pupillary block
- -No increased IOP (15 today)
- -Optic nerve appeared to be grossly intact
- Start atropine (left eye twice daily) in attempt to break synechiae

- 3. Anisocoria (right>left) secondary to posterior synechiae
- 4. Juvenile cataract of left eye
- 5. Anisometropic Hyperopia (left>right)
- -(+)amblyogenic
- Partial Spec Rx
- 6. Amblyopia of left eye
- -Deprivation from cataract v. Anisometropic
- -Re-evaluate after glasses use and with decreased inflammation
- 7. Suppression of binocular vision

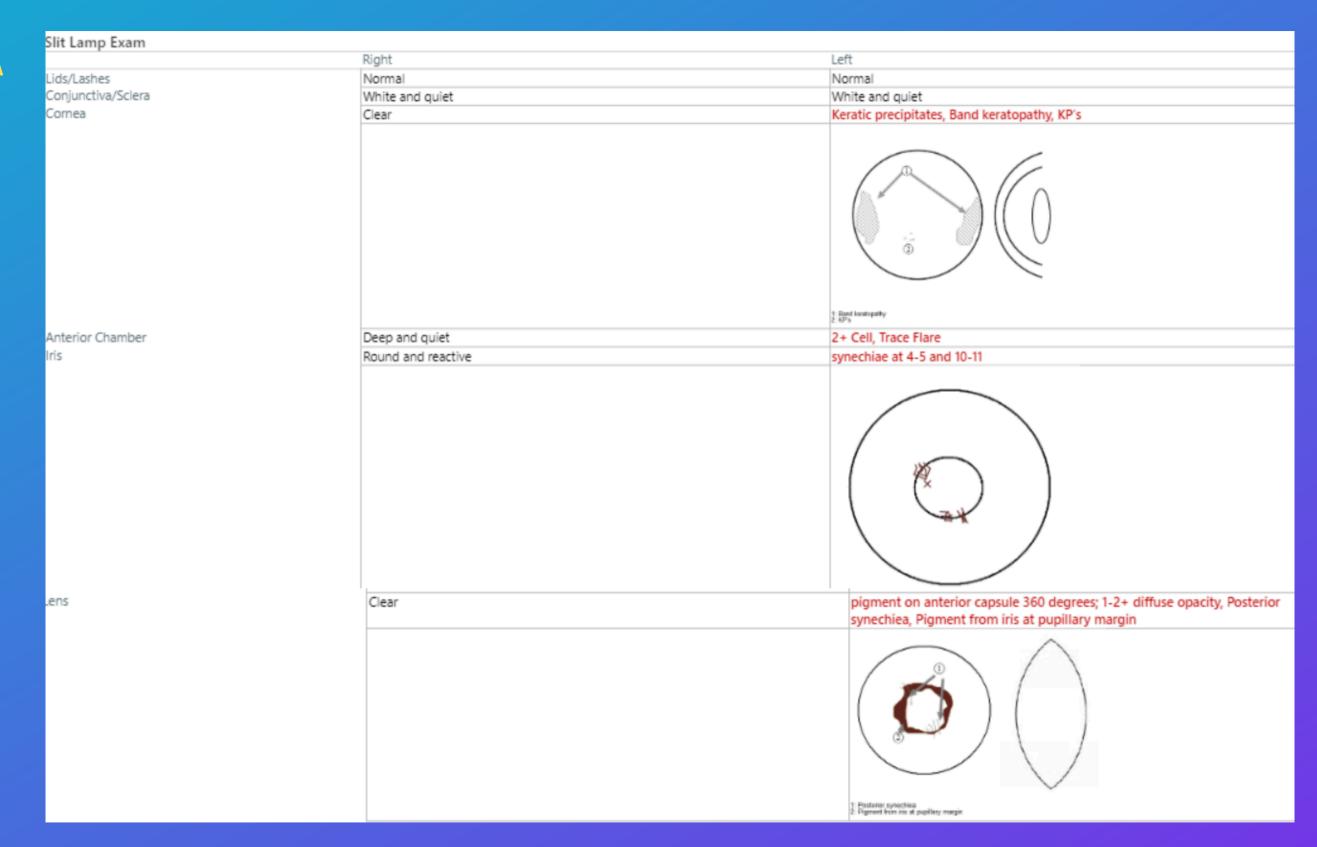
Parent education; RTC in 10-14 days for uveitis check, sooner as needed

LAB RESULTS: POSITIVE ANA

Visit with Pediatric Rheumatologist: Offered Methotrexate pending recalcitrance of uveitis

Visit #2: 2 weeks

- Mom did not fill glasses
- PA and Atropine were being done
- PS were broken
- Flare decreased
- PA decreased to T.I.D.
- Referral made to Pediatric OMD for cataract evaluation



Final Diagnosis & Treatment