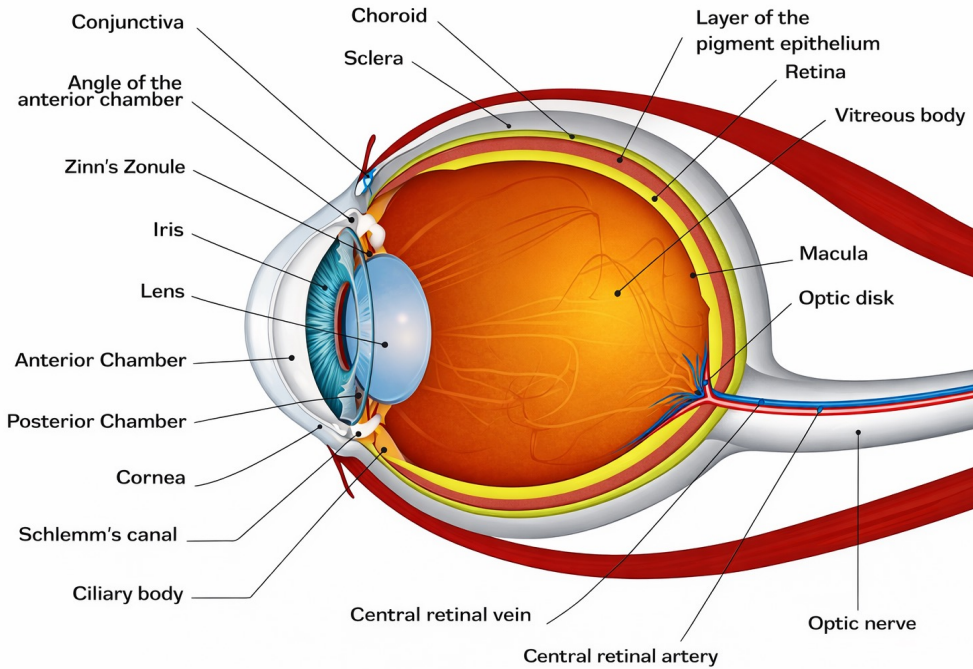


OPHTHALMOLOGY

Basic anatomy and embryology

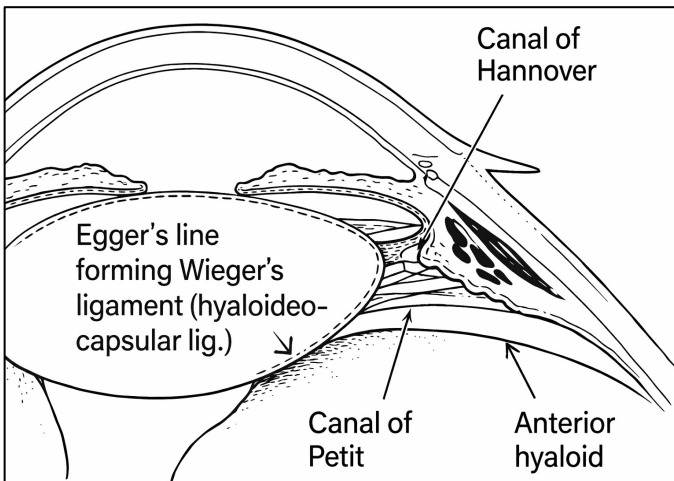


Lens
Epithelium-Cornea, conjunctiva
Lacrimal gland

ON
Retina
Iris and CB epithelium
Sphincter, dilator pupillae

Corneal stroma, endothelium
Sclera (Except superotemporal)
Trabecular meshwork
Iris, choroid stroma
Melanocytes
Ciliaris

Vitreous

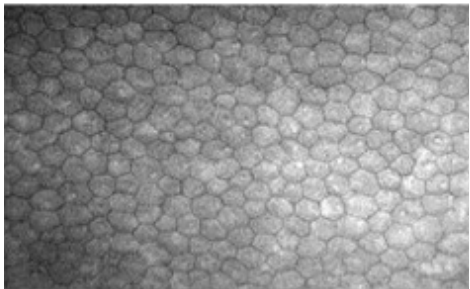
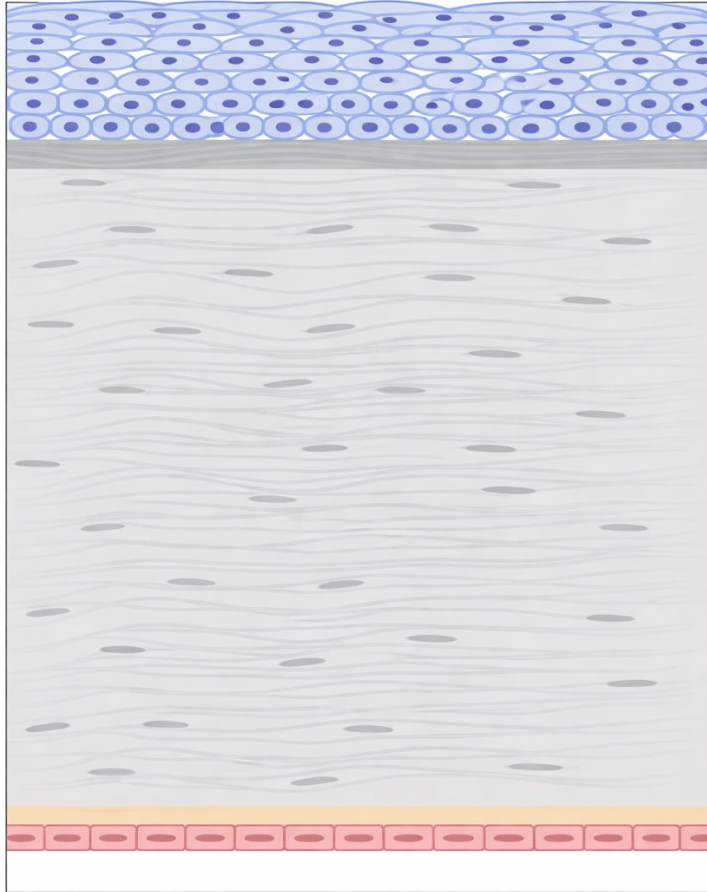


Lamina Fusca
Sclera proper
Episclera
Tenon capsule
Conjunctiva

Retrobulbar:
Peribulbar:

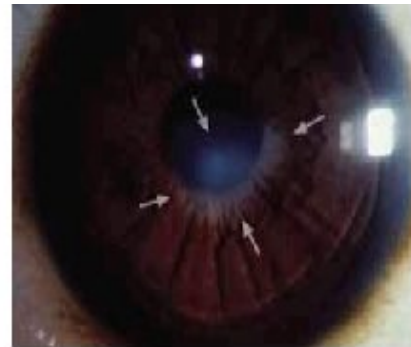
Axial length:
AC depth:
Anisometropia:
At birth:
Total Refractive power:
Cornea
Purkinje images

Cornea



Avascular
Corneal blink reflex:
Corneal Anesthesia:

Interstitial keratitis:



Nebula

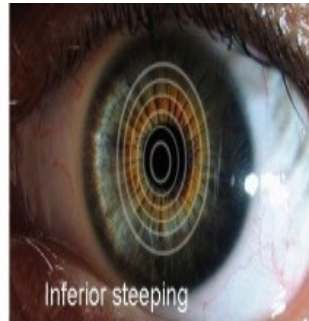
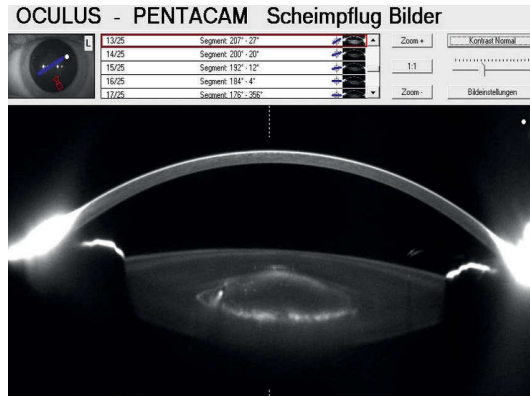


Macula

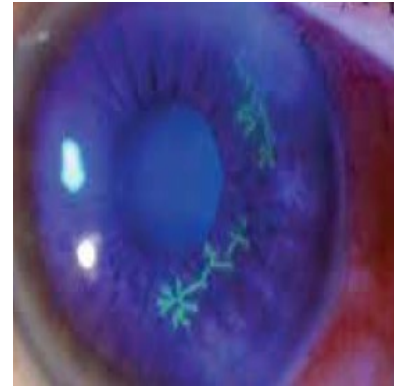
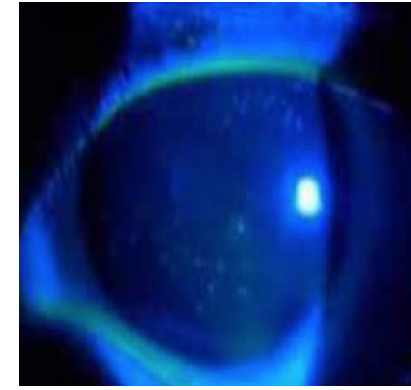
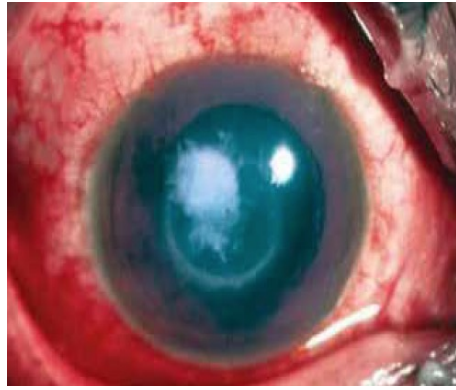


Leucoma

Cornea Investigations



Cornea Infections



Hypopyon:

Sterile, mobile-

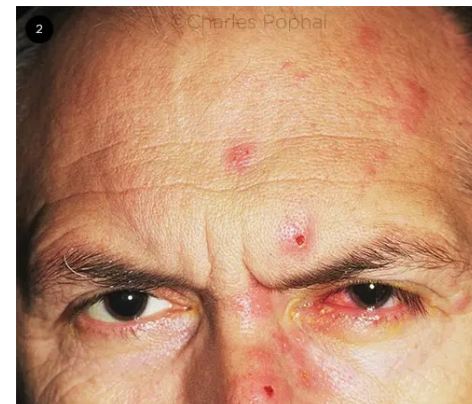
Asterile, immobile-

Ulcus serpens:

- **Wesley's immune ring**
- **Satellite lesions**
- **h/o trauma with vegetative matter**

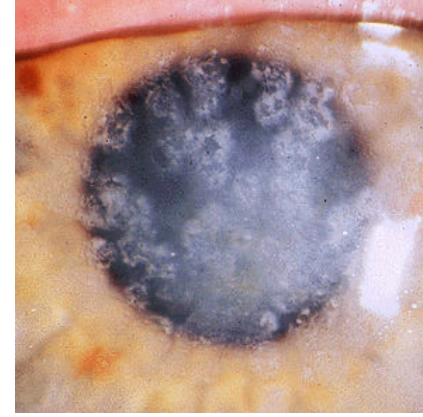
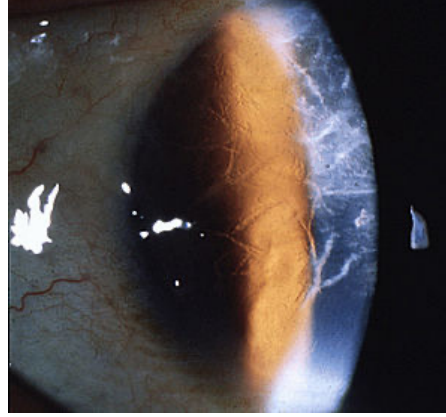
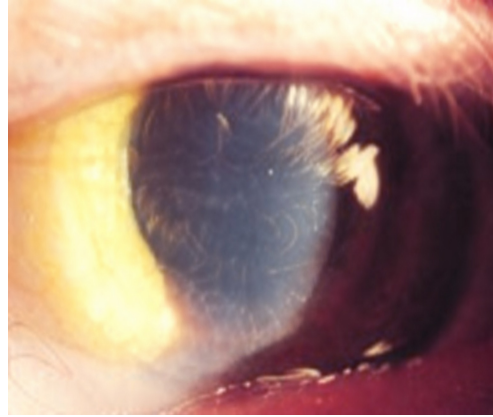
- **Ring abscess**
- **-Pseudo-dendrites**
- **H/o contact lens misuse**
- **PAIN!!**
- **Stain:**

Stromal / disciform keratitis

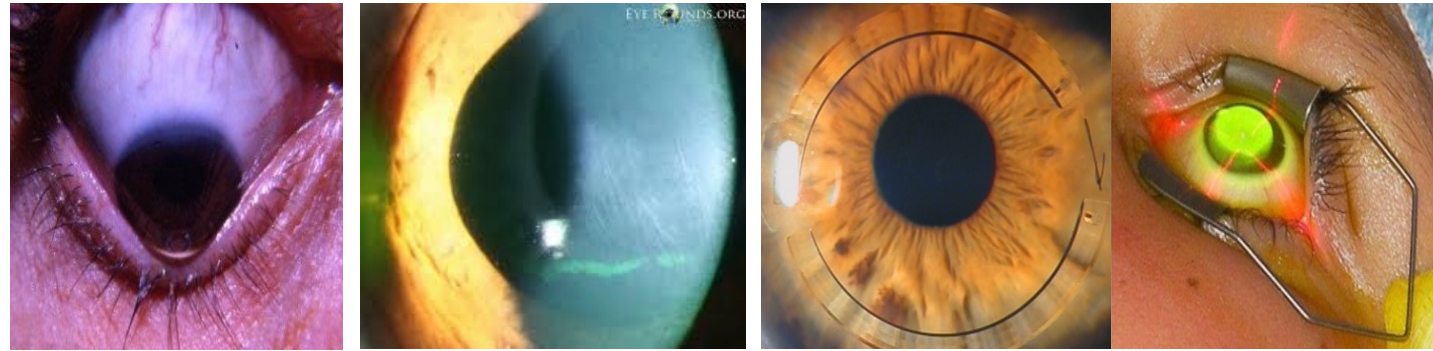


Corneal Dystrophies

Inflammatory?
Progressive?
B/L?
Inherited?

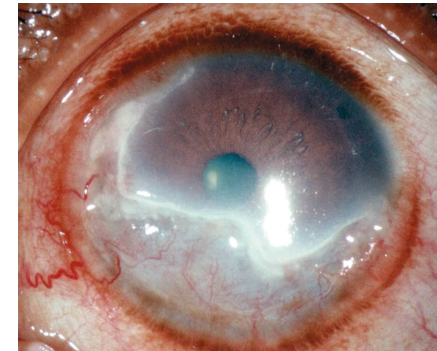


Other corneal-scleral pathologies

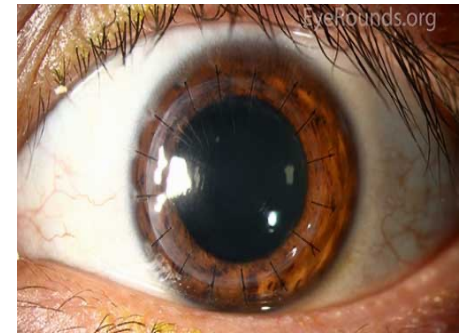
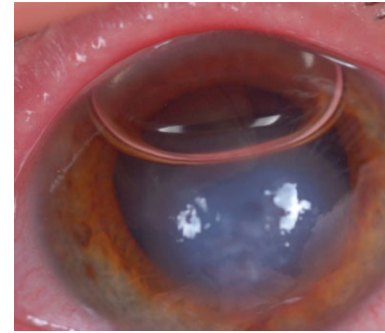
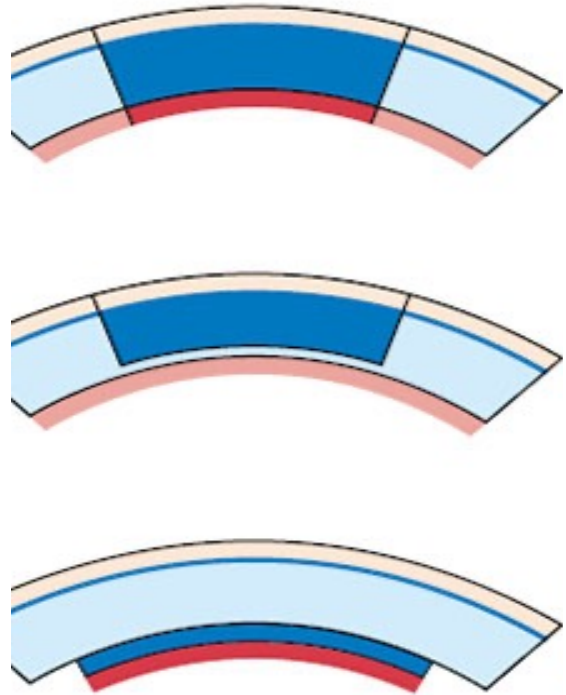


Pain:
2.5% phenylephrine:
Association:

Irregular myopic astigmatism
Munson sign
Vogt's striae
Fleischer ring
Pseudo-Fleischer ring:
Rizutti sign
Oil droplet ("Charleaux" sign):
Scissoring:

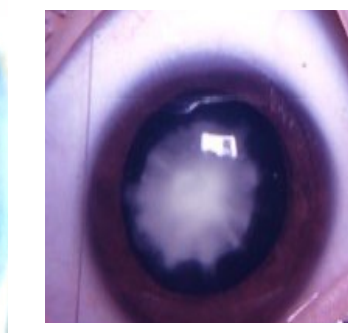
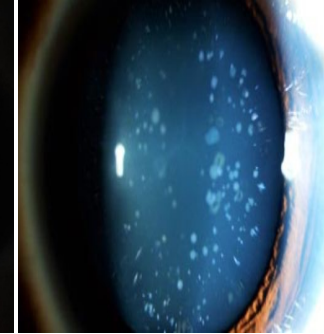
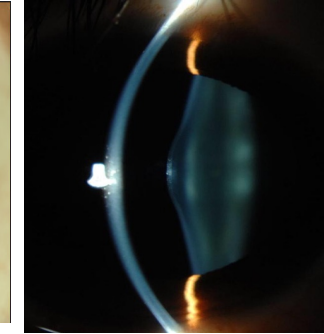
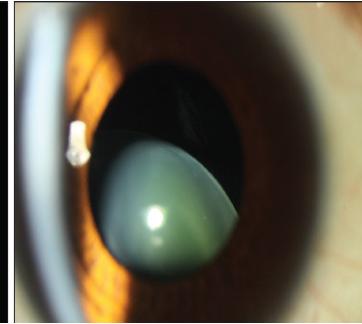
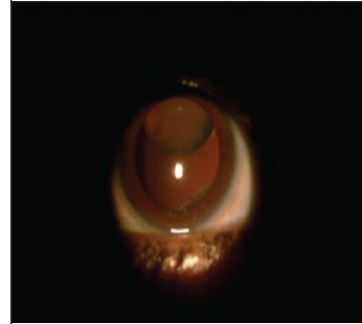
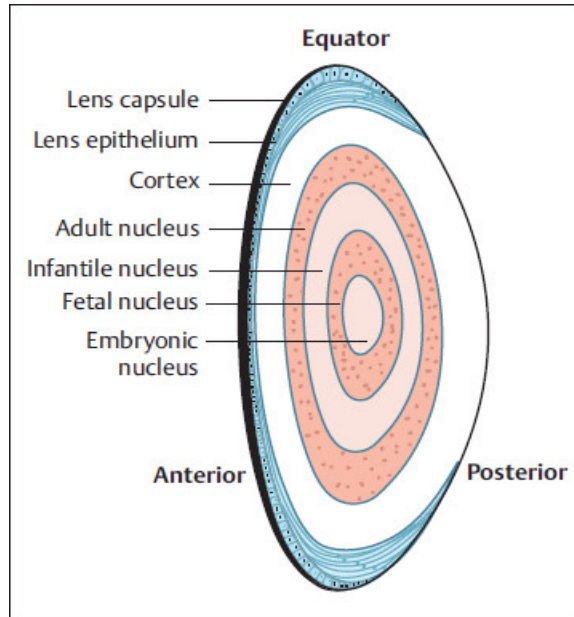


Keratoplasty



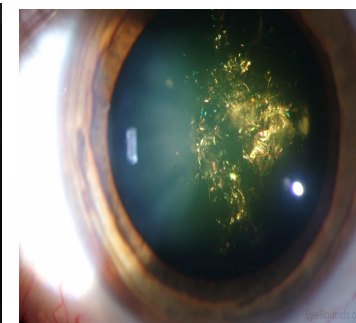
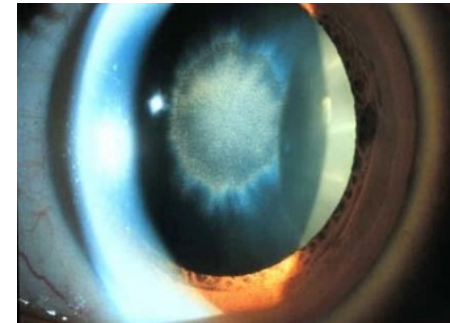
McCarey Kauffman medium
Nylon sutures
Kayes dots: Subepithelial
Khoda-daust line

Lens



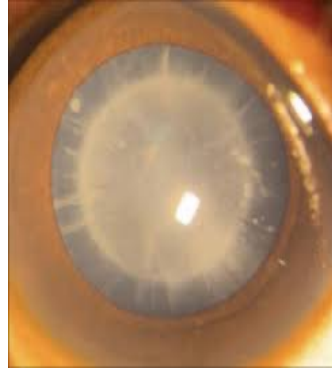
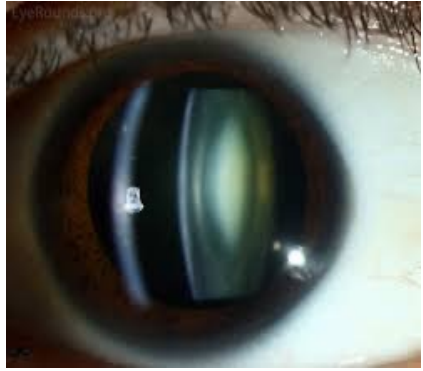
**Superotemporal:
Inferonasal:
Microspherophakia:**

**Patellar fossa
Avascular
Refractive index-1.41 - 1.38-
Water soluble = crystallins
MC: Largest:
HM1,2 – Water soluble**



Fluctuating myopic shifts:

Cataract

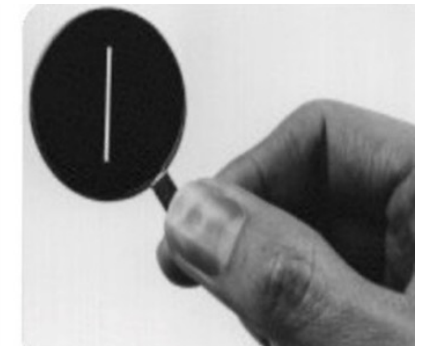


Second sight
LOV Day>night

Phacomorphic glaucoma
Phacolytic glaucoma
Phacotoxic glaucoma
Phacoanaphylactic uveitis

Polychromatic lustre / Breadcrumb:
MCC:
Glassblower's cataract:
Steroids NF2
Shield-like:
Amiodarone Chlorpromazine Au

Difficulty in bright light
Glare

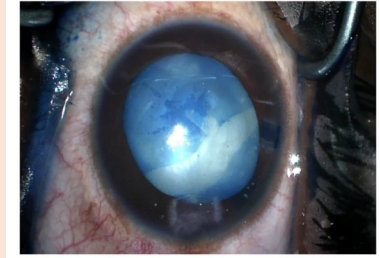


Cataract surgery

ECCE:
Conventional-
SICS-
Phacoemulsification-
MICS-
FLACS-

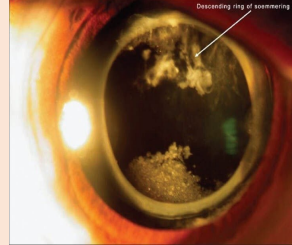
Intra-op:

- Posterior capsular rupture
- Vitreous loss
- UGH
- Expulsive choroidal hemorrhage

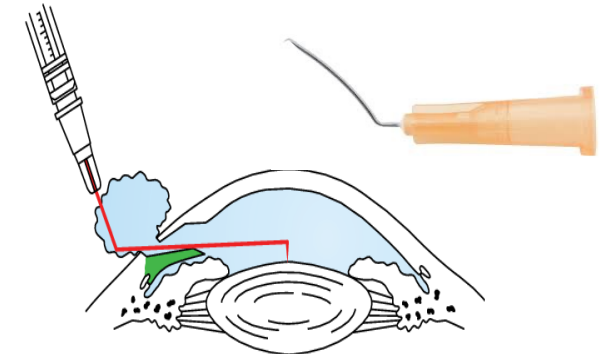
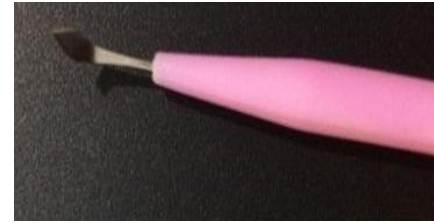


Post-op

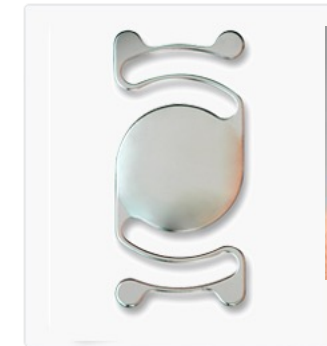
- **PCO**
- **Rx:**
- Endophthalmitis
- TASS
- Irvin Gass syndrome
- IOL displacement
- Endophthalmitis:
 - **MCC early onset-** **Late onset(>6weeks)-**
 - **Rx:**



Cataract Instruments

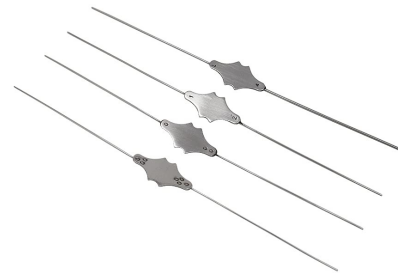
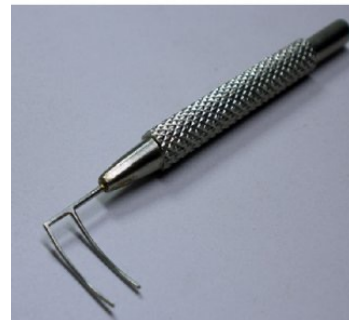


Trypan blue:



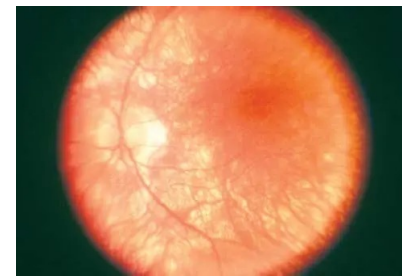
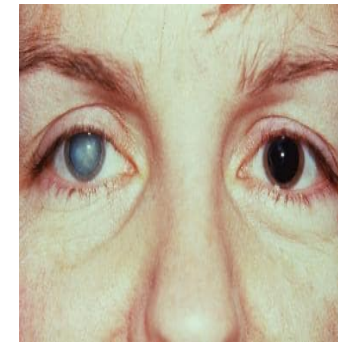
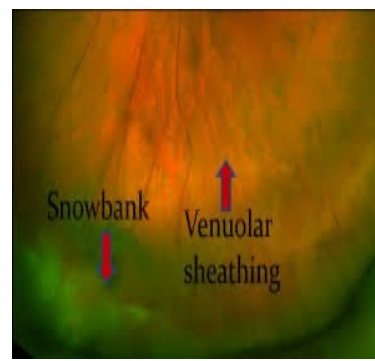
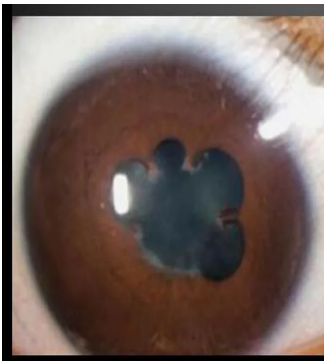
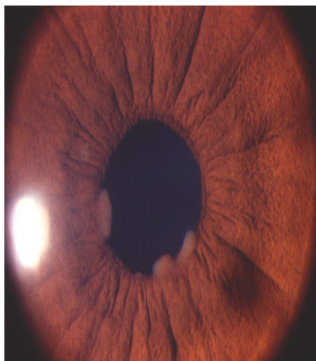
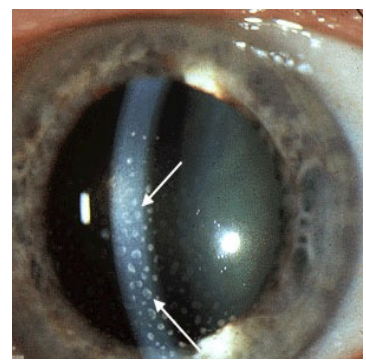
Biometry: Corneal curvature + Axial length
SRKT
Holladay / Hoffer Q
Myopic LASER: Haigis L

Instruments



Uveitis

Iris+ pars plicata
Management:
Pars planitis
Chorio-retinitis

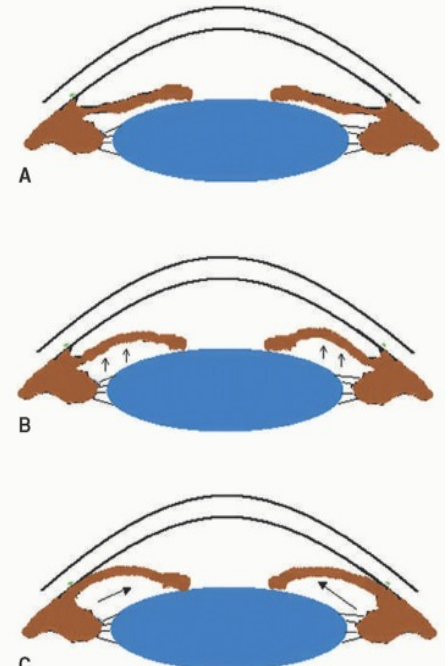
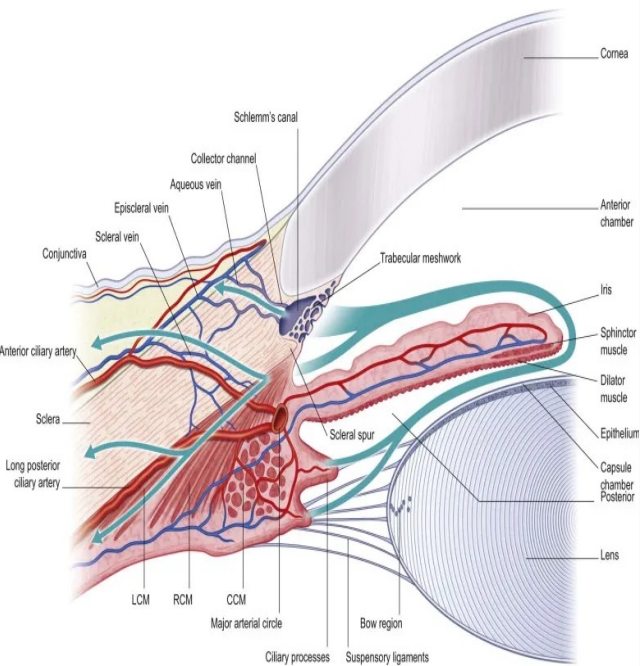


Mutton fat-
Fine granular-
Pigmented-
Stellate-

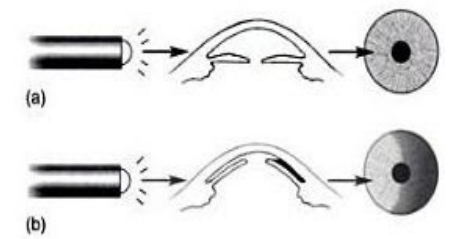
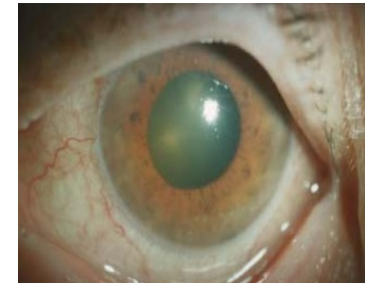
Amsler sign:

Suiguira sign

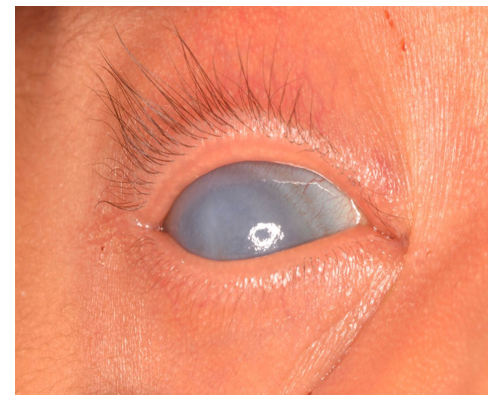
Glaucoma



Angle Closure Glaucoma
 AC: (vs OAG:)
Triad:
 Glaucomflecken (anterior subcapsular opacity)
 Iris atrophy
 Pigment dispersion on endothelium
Management of ACG:
 A
 M
 P
 Surgery:
 POAG:

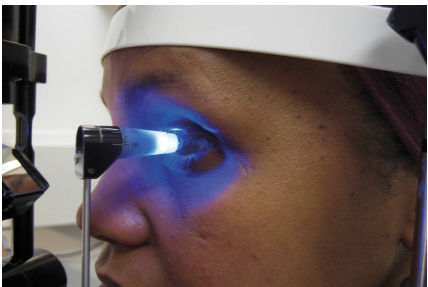
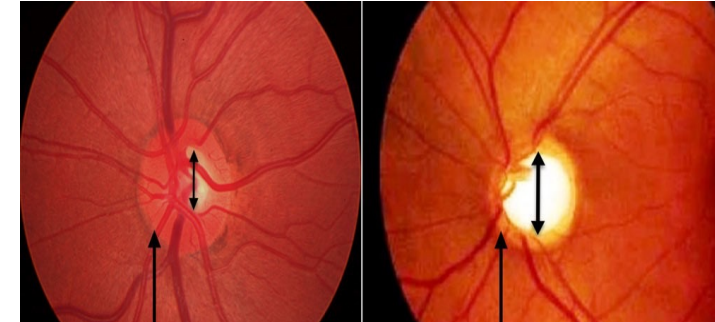
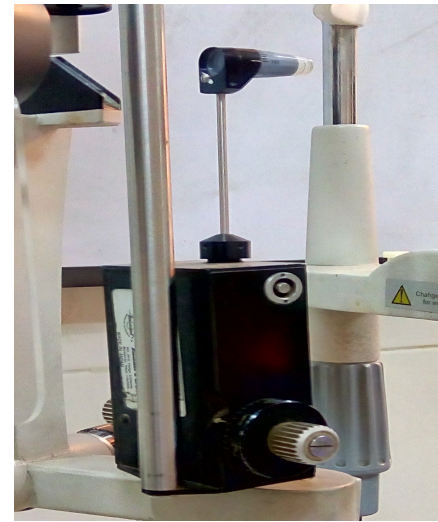


- Unilateral IOP spikes + uveitis
 - Malignant glaucoma/ ciliary block/ aqueous misdirection syndrome: Shallow AC
- DOC:**
- Krukenberg spindle



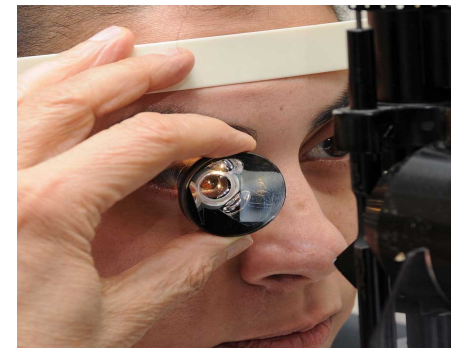
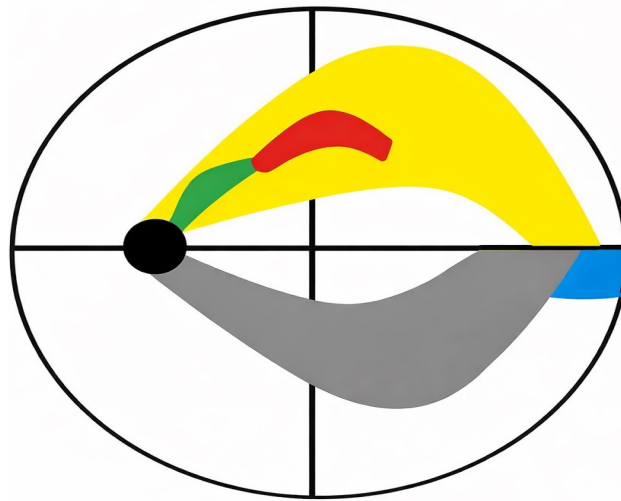
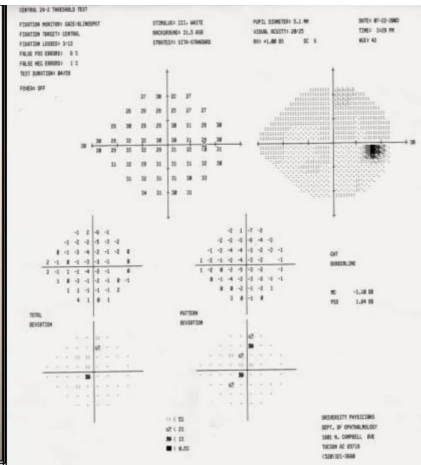
Blepharospasm
Photophobia
Lacrimation
Striae:
 Corneal diameter > 13mm
 Rx:

Glaucoma - Investigations



Mackay-Marg tonometer:

- Increased cup:disc ratio
- Thinning of neuro-retinal rim
- Shifting of vessels
- Laminar dot sign
- Bayonetting sign



Humphrey/ Octopus/ Oculus:

Goldmann perimeter, Bjerrum, Lister:

Visual field of Left Eye

- Nasal step
- Paracentral scotoma
- Seidel scotoma
- Arcuate scotoma
- Biarcuate scotoma

Total internal reflection

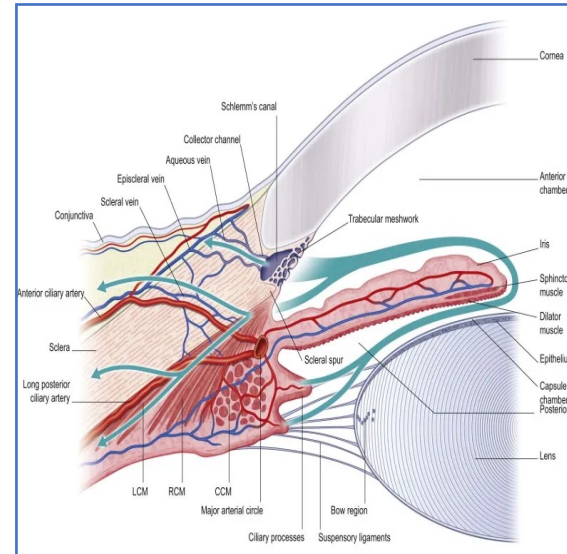
“I can see till Schwalbe's line”

Richardson/ Koeppe:

Zeiss/ Goldmann:

**Earliest-
Paracentral scotoma
Siedel's scotoma
Bjerrum/ Arcute scotoma
Ring/Double arcuate
Roenne's nasal step
Loss of central vision
Temporal crescent loss**

Side	Range
Superior	50°-60°
Inferior	70°-80°
Nasal	60°
Temporal	100°-110°



Anti-Glaucoma Drugs

Decrease Aqueous Production

Ocular Beta Blockers

1. Timolol
2. Betaxolol
3. Levobunolol
4. Carteolol

Nonselective Alpha Agonist

Dipivefrine

Alpha-2 Agonist

1. Apraclonidine
2. Brimonidine

Carbonic Anhydrase Inhibitors

1. Acetazolamide
2. Brinzolamide
3. Dorzolamide

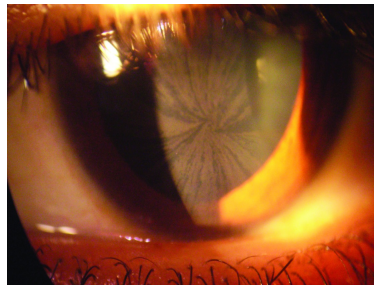
Increase Trabecular Outflow

Cholinomimetic Drugs

Pilocarpine

Rho kinase inhibitor

Netrasudil



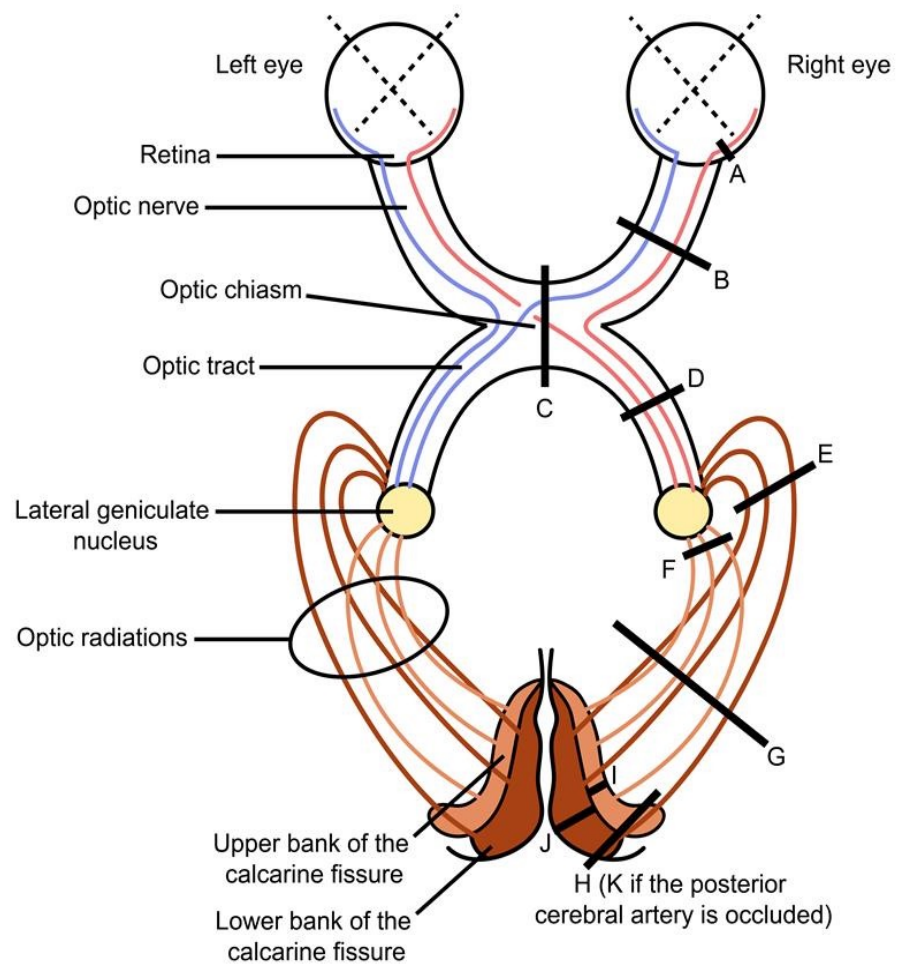
Increase Uveoscleral Outflow

PGF_{2α} Analogues

1. Lantanoprost
2. Bimatoprost

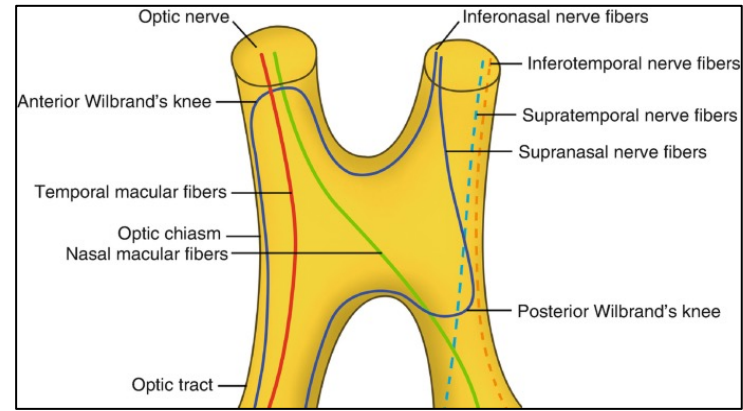
Omidenepag

Visual Field Defects



Visual field deficit

A) Central scotoma		
B) Monocular vision loss		
C) Bitemporal hemianopia		
D, G, & H) Contralateral homonymous hemianopia		
E & J) Contralateral superior quadrantanopia		
F & I) Contralateral inferior quadrantanopia		
K) Contralateral homonymous hemianopia with macular sparing		

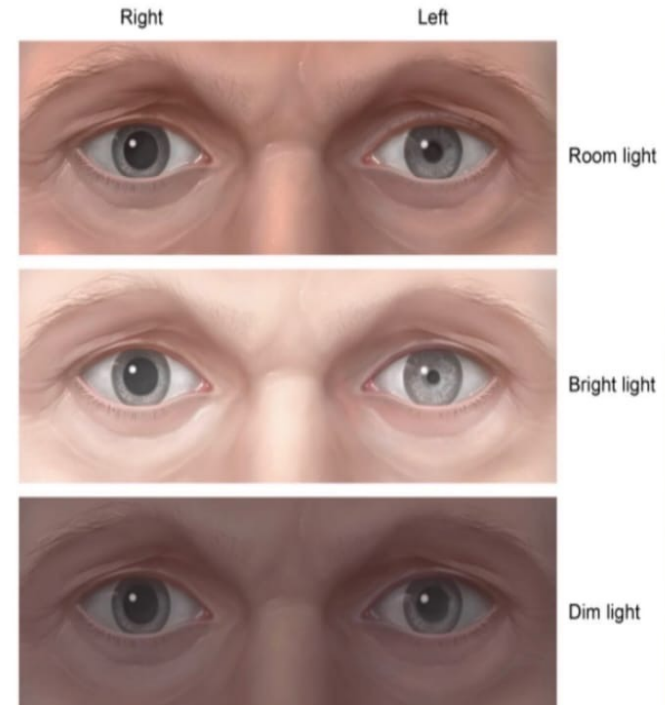
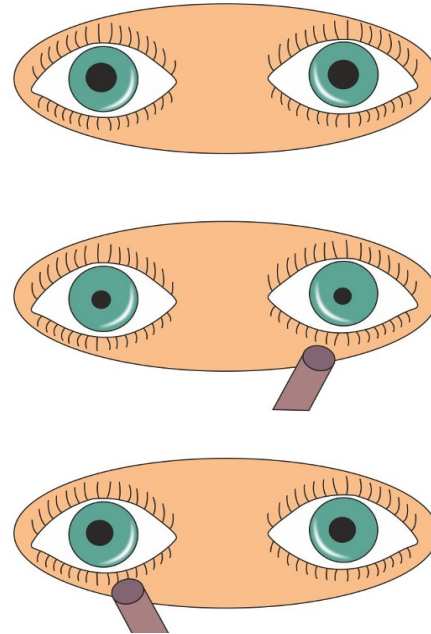
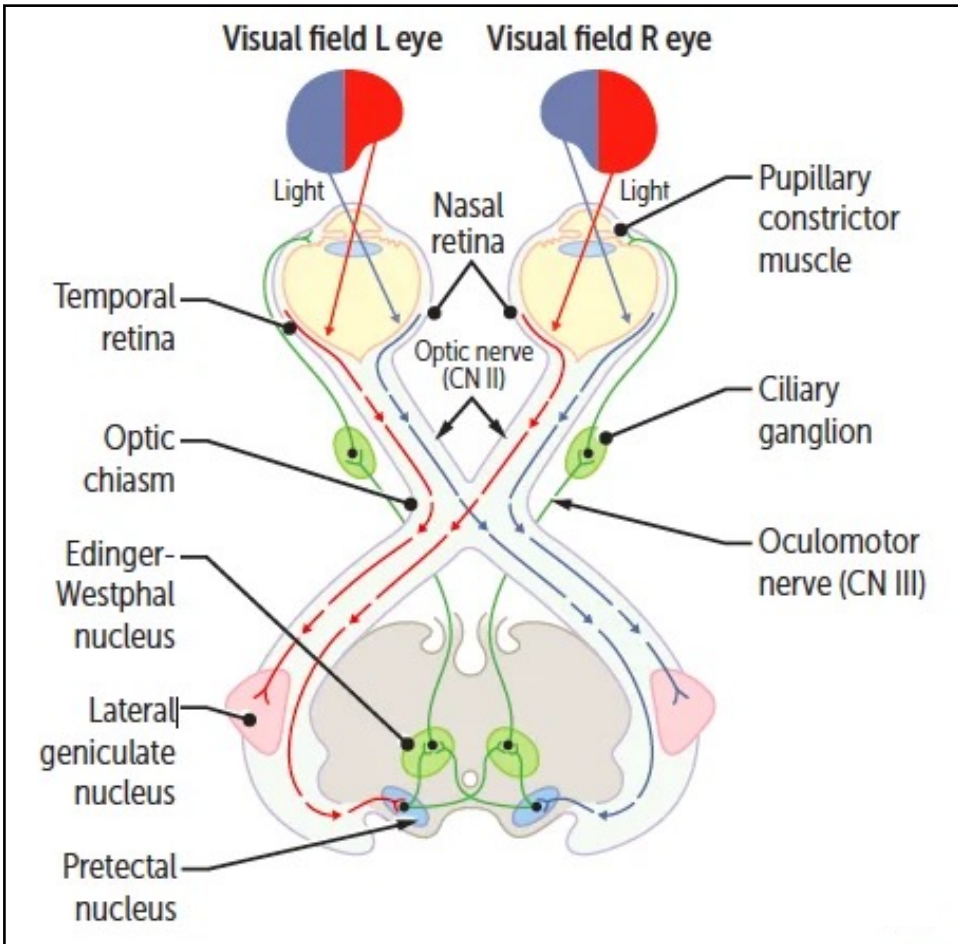


Wernicke's hemi-anopic pupil

Keyhole defect

Light Reflex

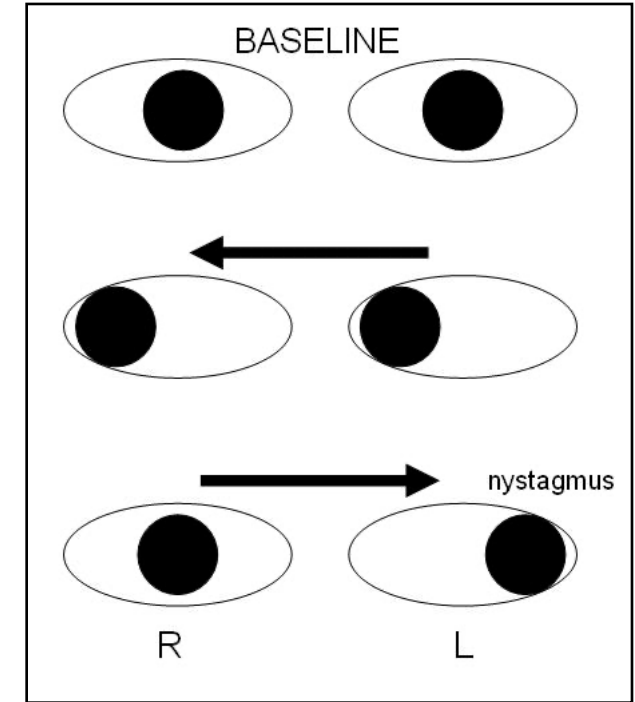
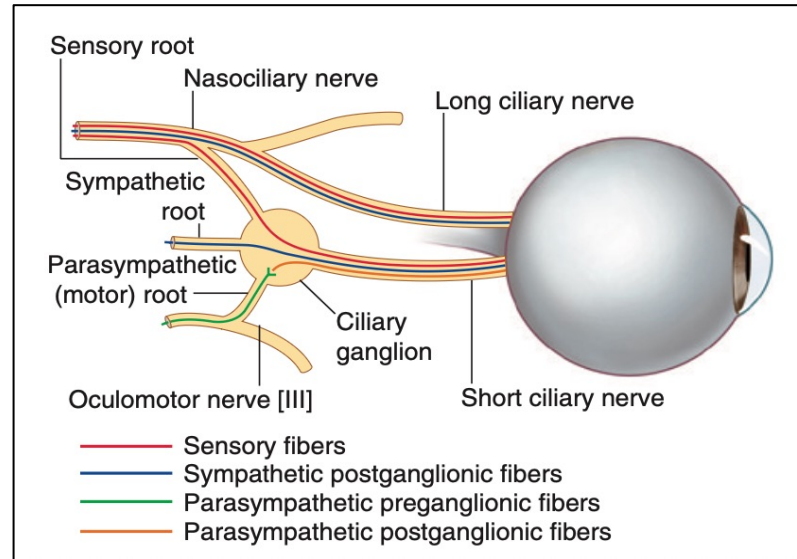
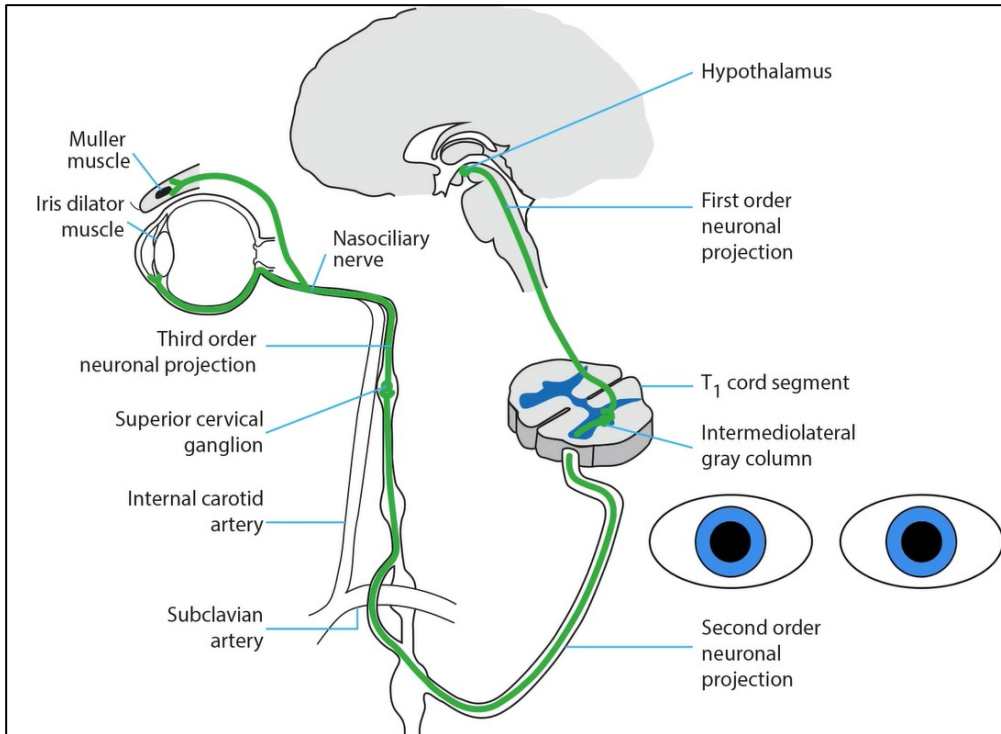
X



	Adie's Tonic pupil	Pharmacological mydriasis
I/L Dilated pupil		
Accommodation		
0.125% Pilocarpine		
1% Pilocarpine		

Sympathetic pathway

INO



4% Cocaine test

Mechanism: Blocks NE reuptake

→ No NE in postsynaptic cleft

→ No dilation: Horner's syndrome

1% Apraclonidine test

•Mechanism: Denervation

hypersensitivity of α_1 receptors

•Pupil dilates: Horner's syndrome




1% Amphetamine test

•Mechanism: Releases stored norepinephrine from presynaptic terminal

•Pupil dilates → Pre-ganglionic lesion

•Pupil does NOT dilate → Post-ganglionic lesion

Optic Atrophy

Sign	Primary	Secondary	Consecutive
Disc colour	Chalky white	Grey	Waxy pale
Disc margins	Distinct	Blurred	Attenuated arteries
Cause	<ul style="list-style-type: none"> Optic N compression Hereditary / Nutritional Optic Neuropathy 	<ul style="list-style-type: none"> Chronic papilloedema Papillitis Anterior Ischemic Optic Neuropathy 	<ul style="list-style-type: none"> Chorioretinal disease (e.g. RP) Central Retinal Artery Occlusion
Image			

Multiple sclerosis

- Retrobulbar neuritis
- U/L LOV + No VFD
- Fundus normal
- Pulfrich sign
- Uthoff sign

IOC:
TOC:

Papilledema

Painless B/L
VFD:
Paton's lines

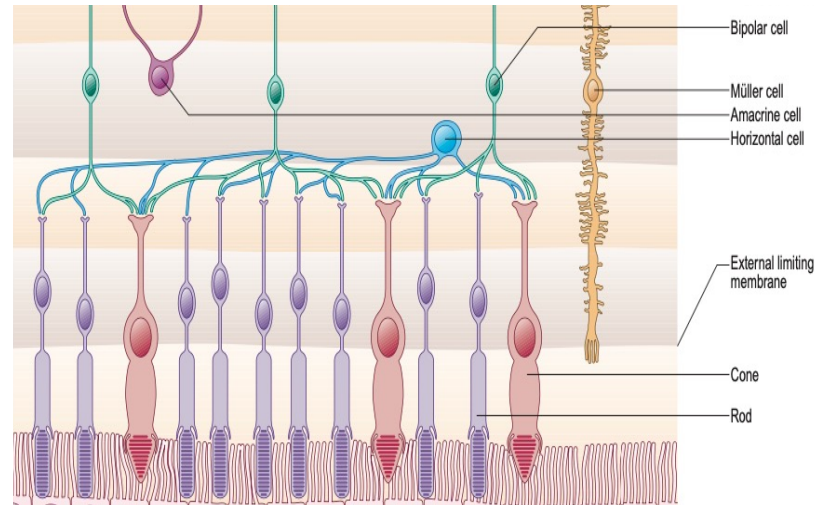
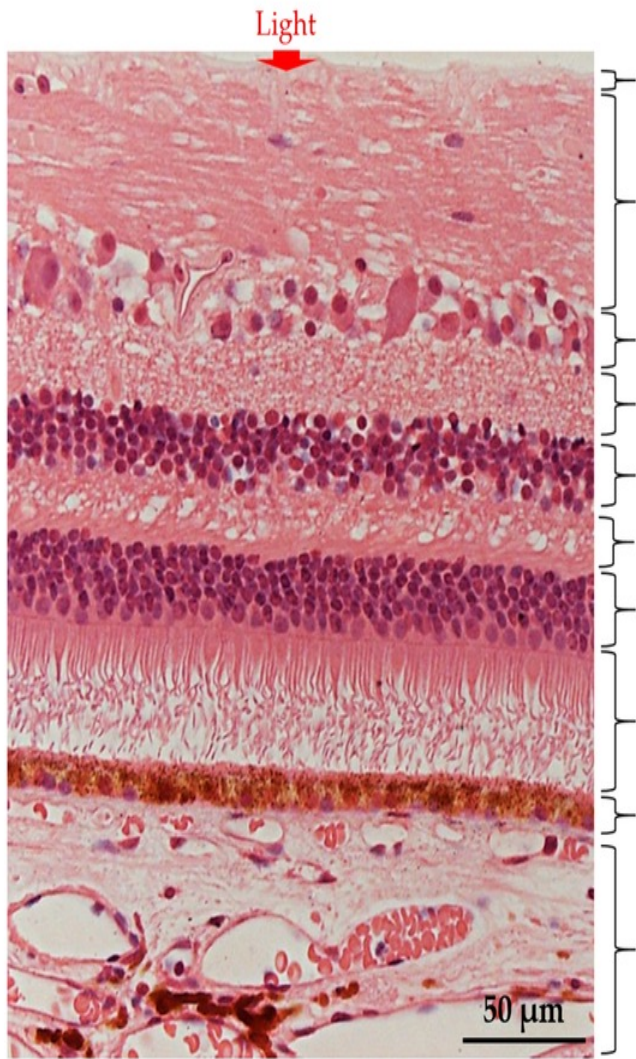
Altitudinal defects:

- Glaucoma
- ON drusen
- AION

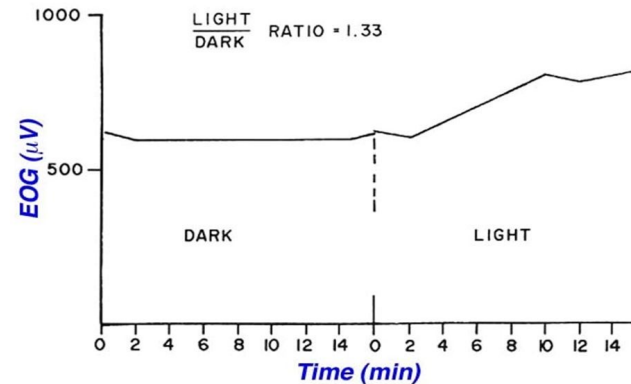
Absent lamina cribrosa
Retrodisplacement of ON



Retina



Light: Hyperpolarisation in rods-
Glutamate release decreases

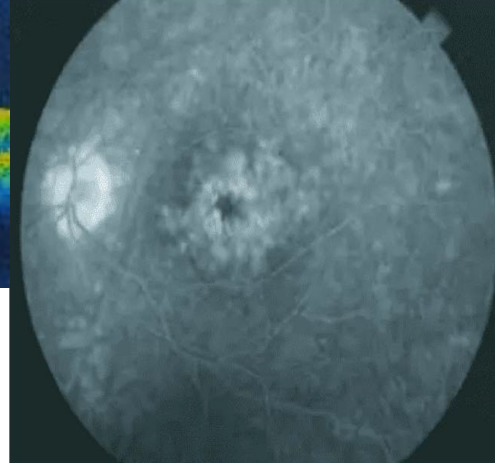
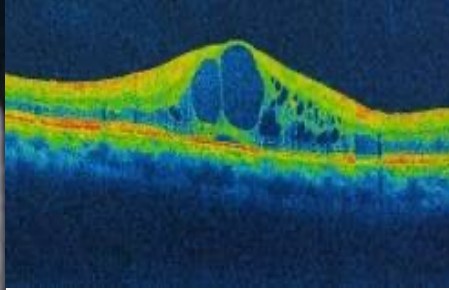
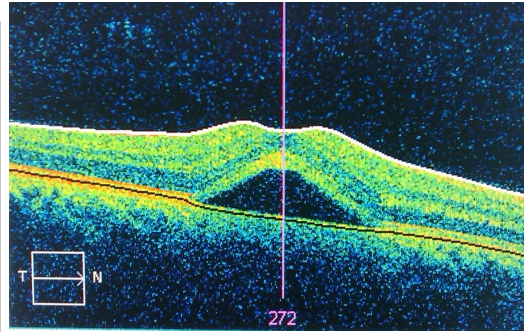
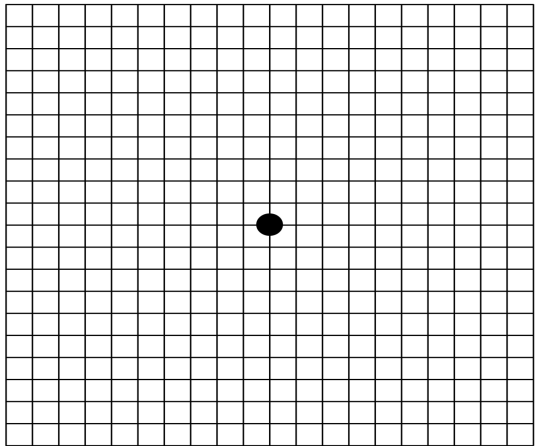


Feature	Direct	Indirect (+20 Lens)
Magnification	15X	3X
Field	2DD	8DD
Orientation	Direct	Inverted
Periphery	Limited	Full
Stereopsis	Absent	Present



Blood supply:
Outer 4:
Inner 6:

Macular Disorders & Investigations

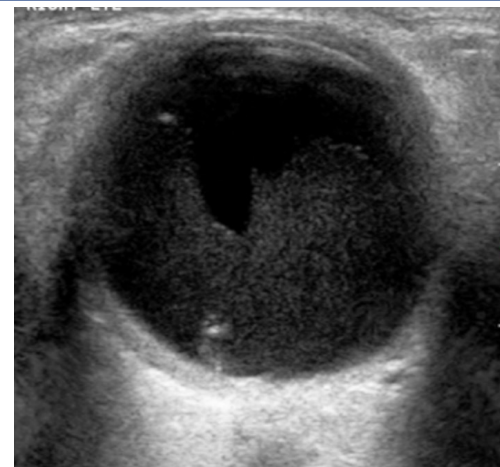
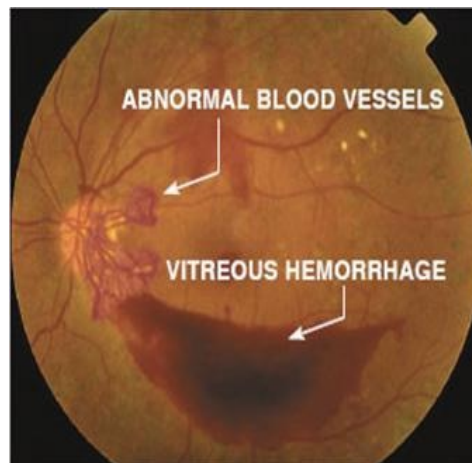
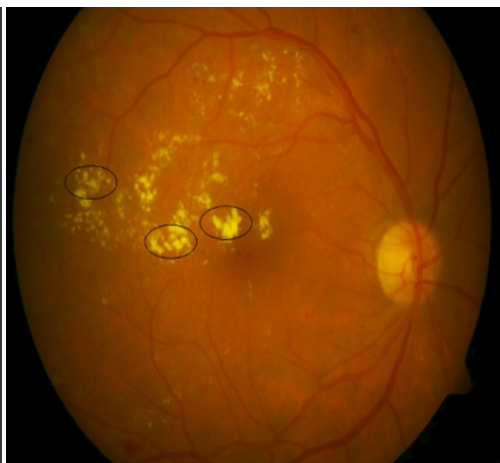
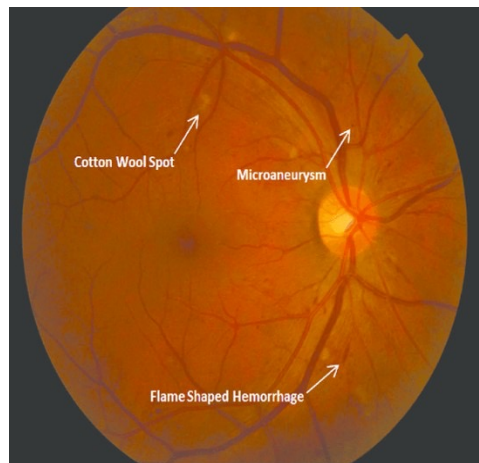


- Diagnosis-
- Layer-
- R/F: Steroids / Type A personality
- Rx:

- Diagnosis-
- Layer-
- R/F- Irvine-Gass
- PG analogue
- Niacin toxicity
- DR
- Uveitis/ RP/ CRVO
- Rx-

- BLOOD RETINAL BARRIER
- Outer : RPE
- Inner: Capillary Permeability

Diabetic Retinopathy



ETDRS
Microaneurysm:
Hemorrhages: Dot and bot:
Hard Exudates:
Flame shaped:
Cotton Wool/ Soft exudates:
Screening-
Type 1
Type 2
Most important R/F:

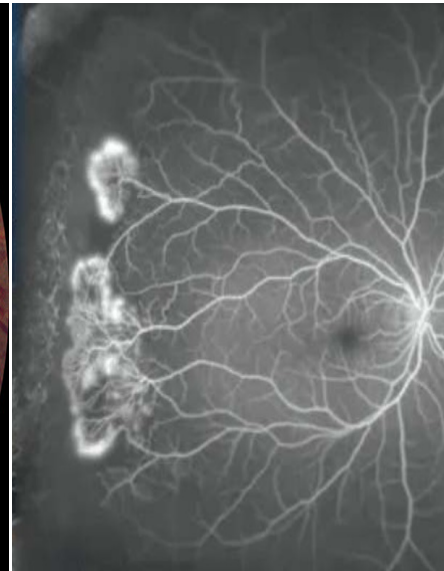
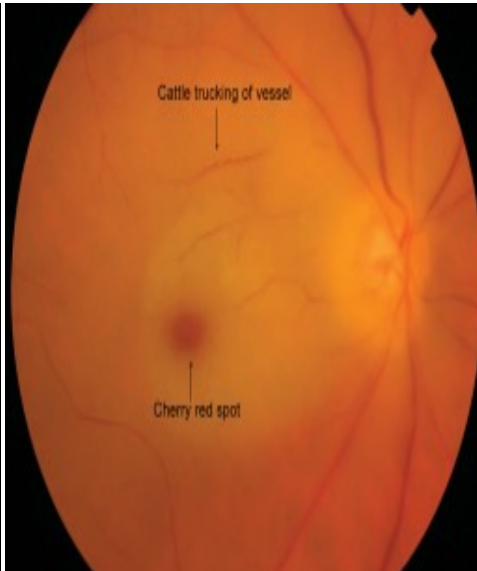
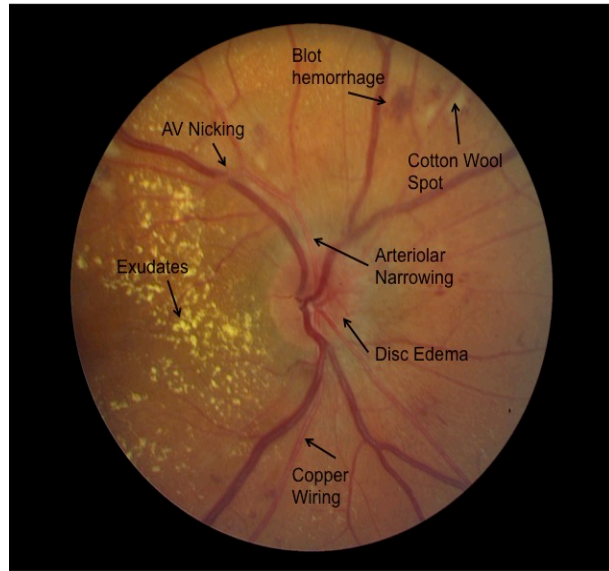
Severe NPDR (4-2-1 rule):

- > 20 intraretinal hemorrhages in each of four quadrants
- Definite venous beading in two or more quadrants
- Prominent IRMA in one or more quadrants

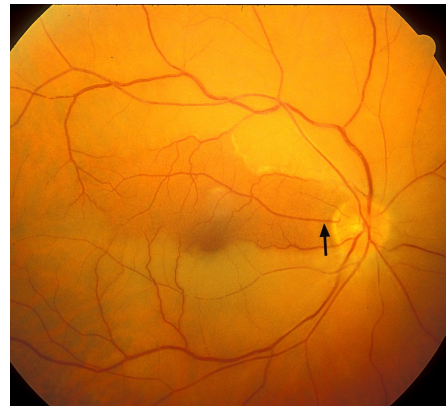
Advanced Diabetic Eye Disease:
Persistent vitreous hemorrhage
Tractional retinal detachment
Neovascular glaucoma

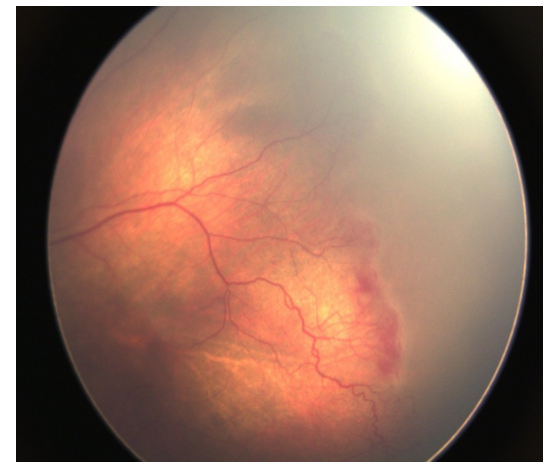
Bevacizumab
Ranibizumab
Brolucizumab
Ramuciruzumab
Aflibercept
Pegaptanib

Retina pathologies



Grade 1	Arteriolar narrowing
Grade 2	Salu sign-S shaped deflection of veins
Grade 3	<ul style="list-style-type: none"> • GUNN: Tapering of vein at either end • BONNET: Dilatation of vein at junction • Flame shaped Hemorrhage • Cotton Wool spots • Copper wiring
Grade 4	Silver wiring Papilledema

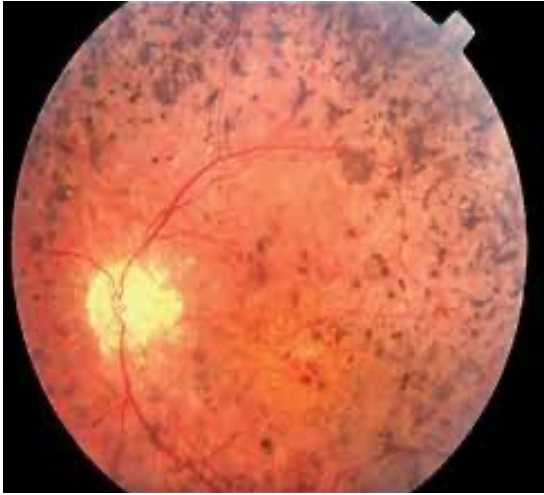




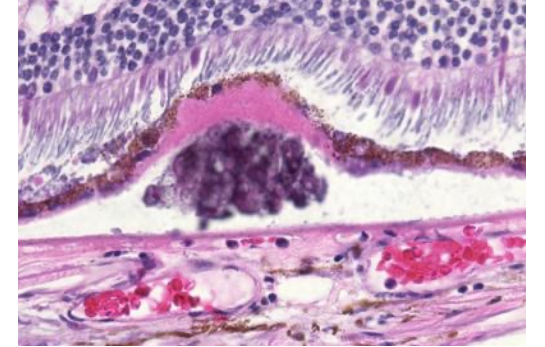
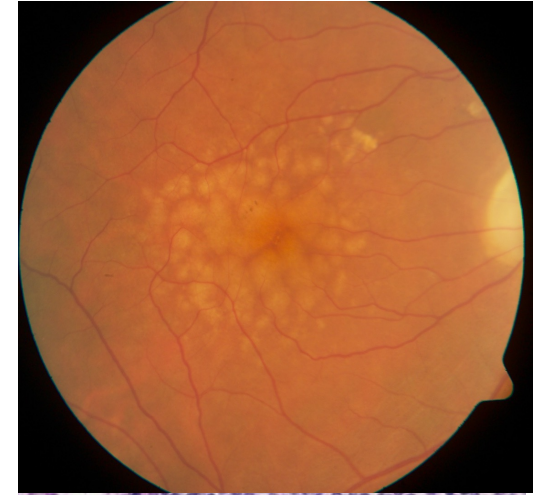
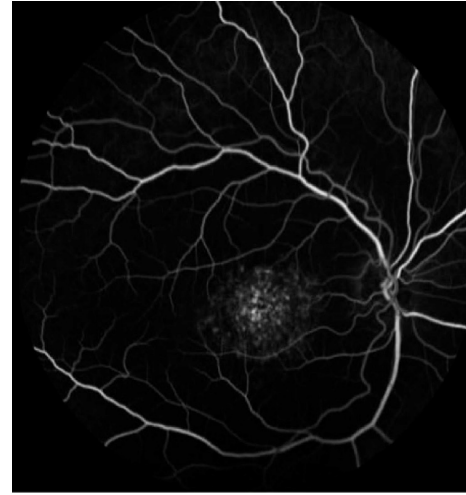
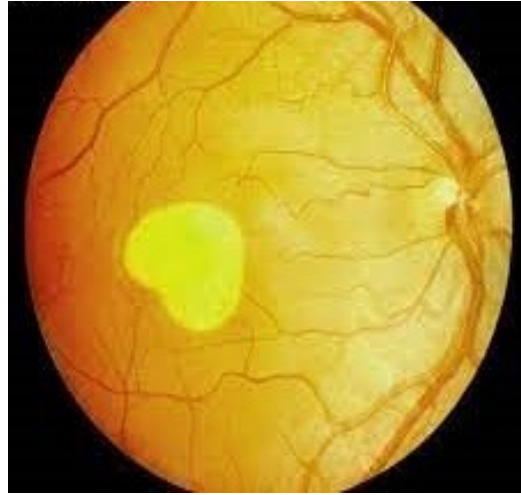
- R/F: Birth weight <1750g
- Gestational age \leq 34 weeks
- Sick infants (oxygen therapy, sepsis)
- Screening:

Stage 1	Demarcation line between vascular and avascular retina
Stage 2	Elevated ridge at the demarcation line
Stage 3	Ridge with extraretinal fibrovascular proliferation
Stage 4	Partial retinal detachment
Stage 5	Total retinal detachment

ROD dystrophy



CONE dystrophy



- MC Inheritance:
- TRIAD:

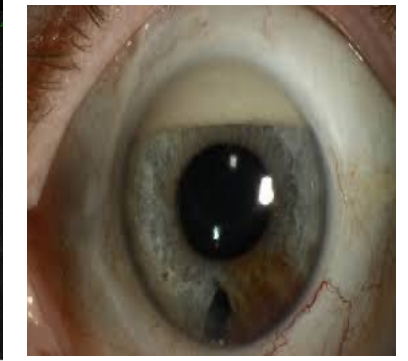
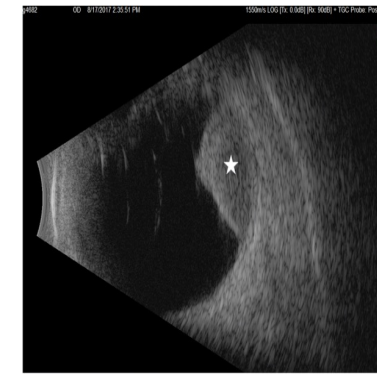
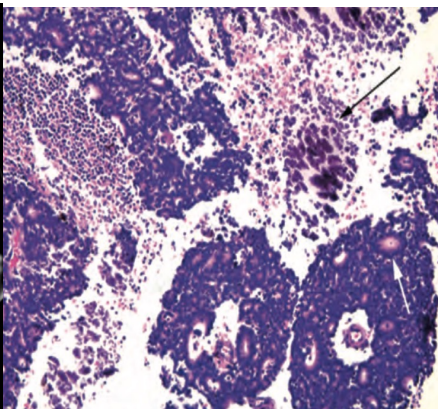
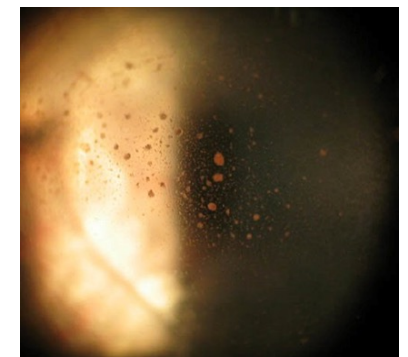
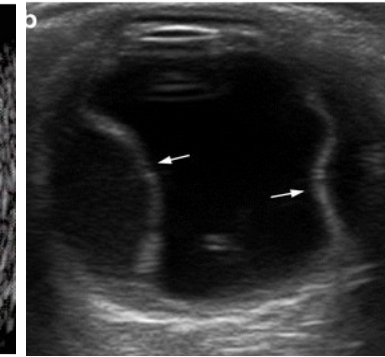
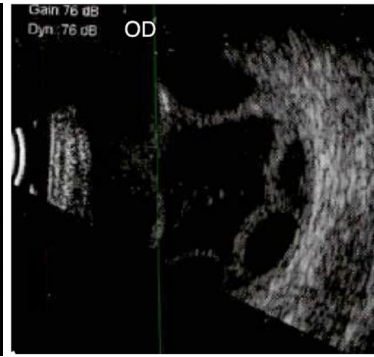
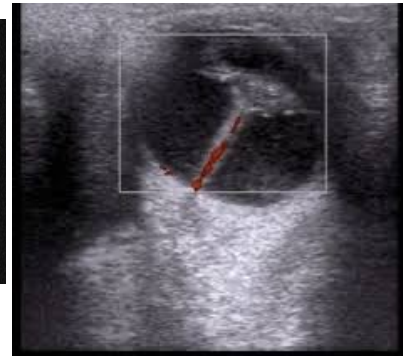
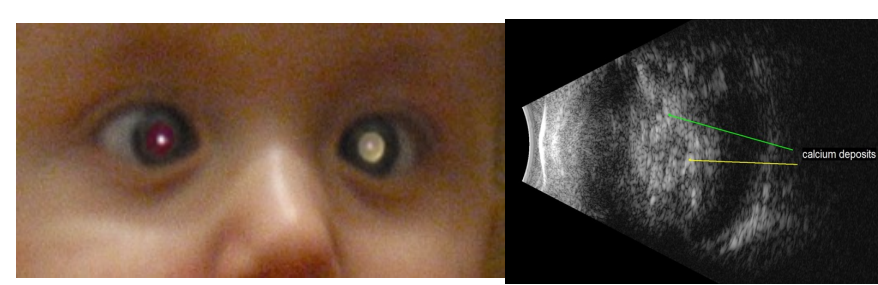
- C/F:
- IOC:
- Docosahexanoic acid ↓
- Usher, Refsum, Kearns-Sayre, Bardet-Biedl

AREDS₂-
Vitamin C, E, Zn, Cu,
Lutein, Zeaxanthin

Oguchi's disease/ congenital stationary night blindness:
Mizuo-nakamura phenomenon

Retinoblastoma

Retinal detachment

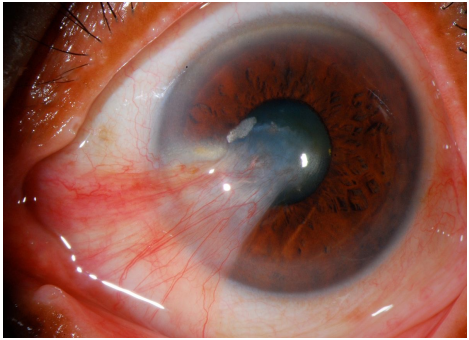


Sporadic:
Gene/ inheritance:
Trilateral RB:
MC association:
C/F-
IOC-
IOC for spread-
TOC-

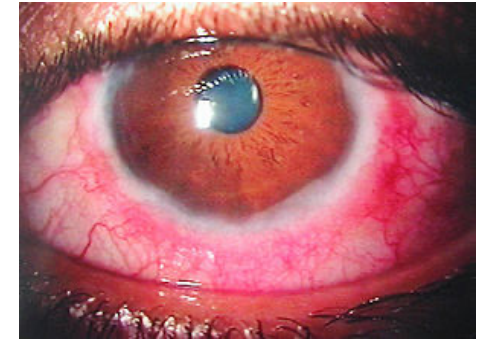
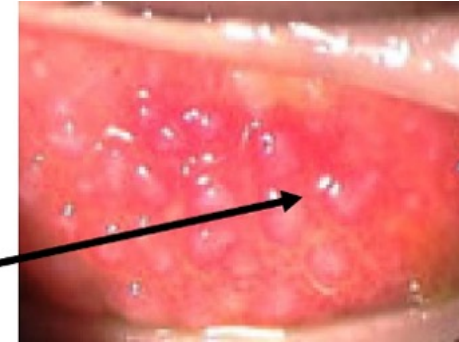
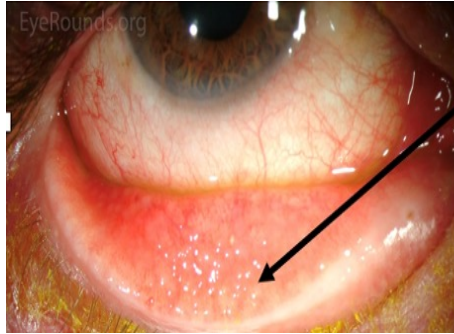
Hereditary:

Rhegmatogenous- Trauma/myopia/surgery
“Curtain falling in front of eye”
Exudative-
Young boys + U/L lipid exudation:
Tractional-
Rx:

Conjunctiva



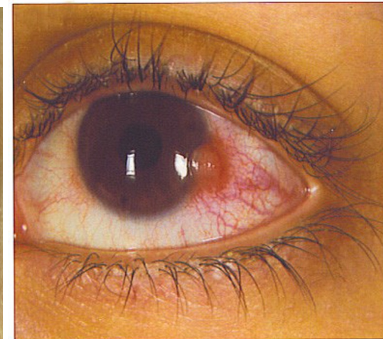
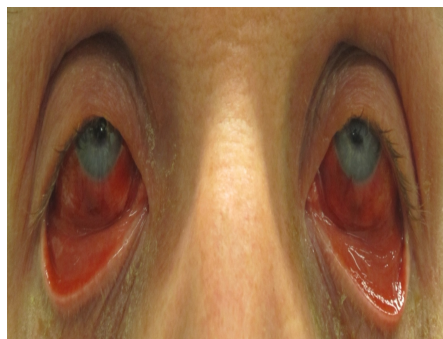
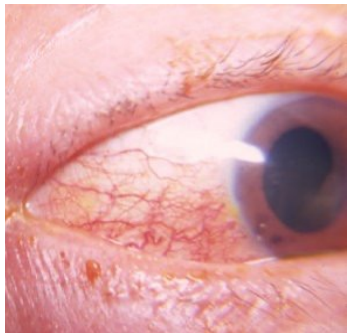
Diagnosis:
C/F:
 Limbus Stem cells
 ABCG2, CD34
Rx:



- Toxins (e.g., Apraclonidine)
- Viruses (e.g. COVID-19)
- Chlamydia

- Bacterial
- Allergic (AKC, VKC)
- Foreign Body
- GPC

- 5-14yr boys
- Ropy discharge
- Hysn:
- Shield ulcer
- Cobblestone papilla
- Horner-Tranta dots
- Pseudogerontoxon
- DOC:



<u>Neonatal conjunctivitis</u>	<u>Onset age</u>	<u>Treatment</u>
Chemical Crede method	<24 hr	Eye lubricant
Gonococcal	24-48hrs	Ceftriaxone
Chlamydial (D-K)	5-14 days	Erythromycin

Trachoma



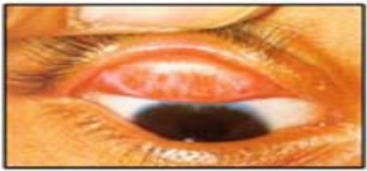
Trachomatous inflammation-follicular (TF)

Five or more follicles of >0.5 mm on the upper tarsal conjunctiva

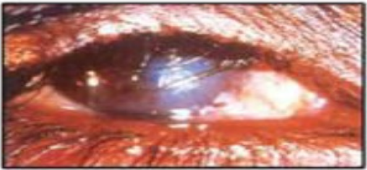


Trachomatous inflammation-follicular and intense (TI)

Inflammatory thickening obscuring more than half of the normal deep tarsal vessels



Trachomatous scarring (TS)



Trachomatous trichiasis (TT)



Corneal Opacity (CO)



Surgery



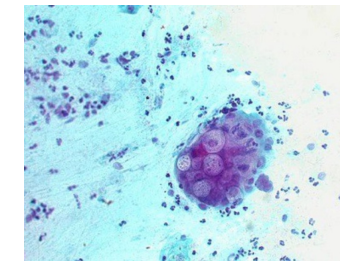
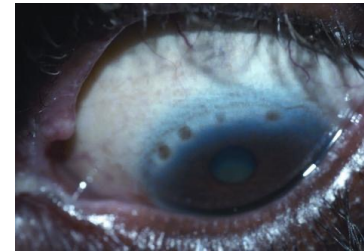
Antibiotics



Facial cleaning



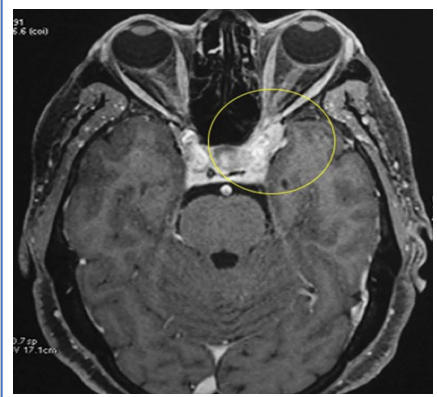
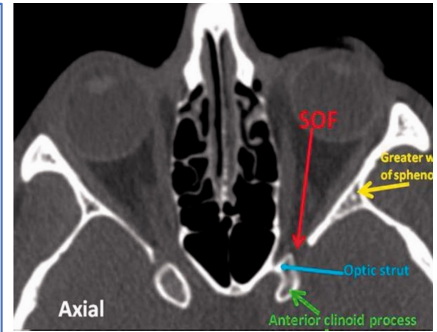
Environmental change



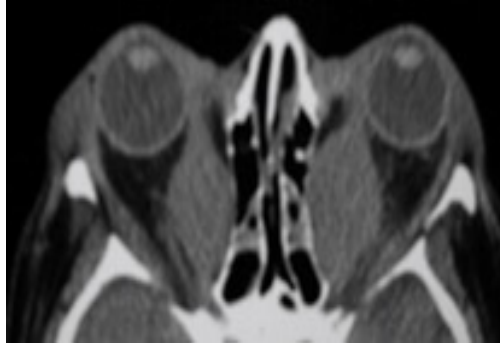
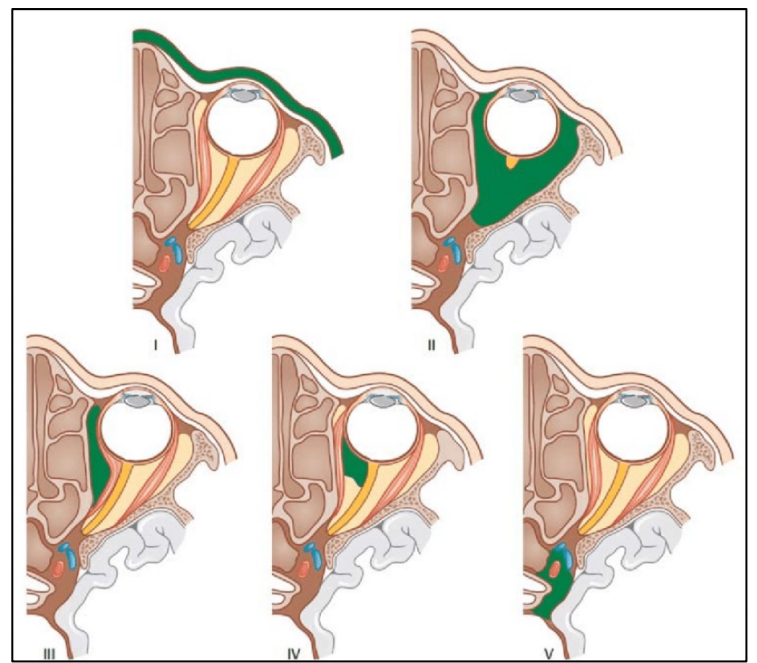
ELIMINATION CRITERIA:
TF of $< 5\%$ in children aged 1 to 9 years
TT $< 0.2\%$ / $< 1/1000$ in adults > 15 yrs

Orbit

Proptosis:
MCC:
Pulsatile:
Increases on bending/ Valsalva:
Increases on crying:
Increases with URTI:
MCC of orbital mets:
Child-
Adult-
MC primary orbital tumor in child
Benign- Malignant-
MC orbital tumor in adult
Benign- Malignant-
MC lacrimal gland neoplasm:



	ORBITAL APEX SX	CST
NERVES		
OPHTHALMOPLEGIA		
VISION LOSS		
U/L OR B/L		
CHEMOSIS, PROPTOSIS		
ONSET		



Lacrimal Apparatus

Watering eye

Syringing and probing

Tear film: Inner-
Middle-
Outer-

Dry eye

Obstruction+

Obstruction-

Hard stop:
Soft stop:

Jones dye test-1

Opacify+

Opacify-

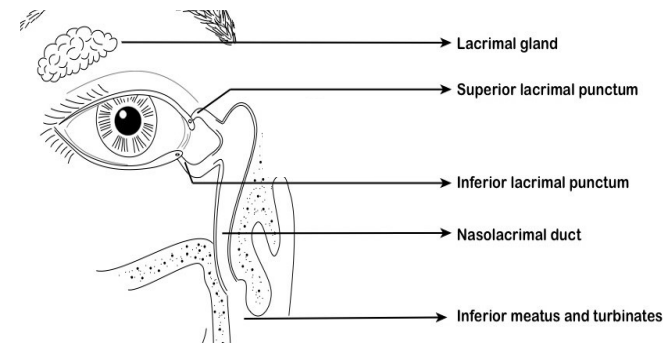
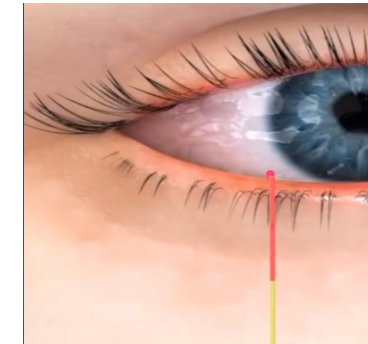
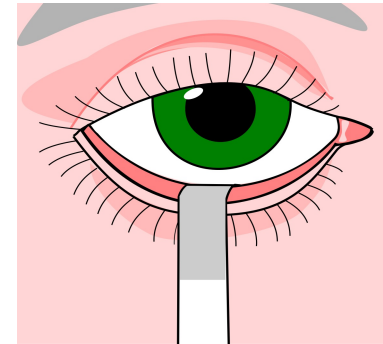
Jones dye
test-2

Opacify+

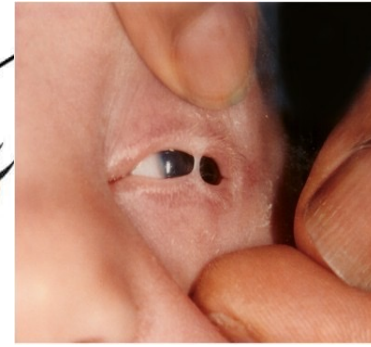
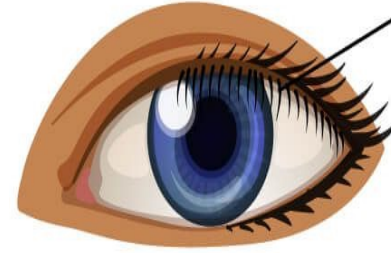
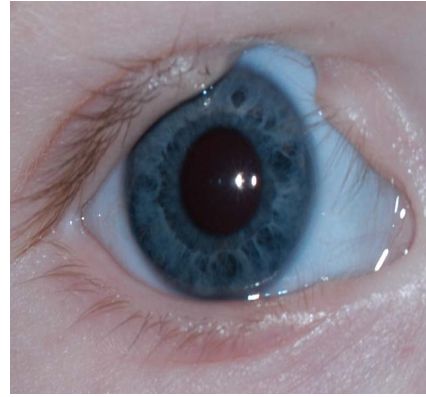
Opacify-

Congenital dacryocystitis: MCC:
Mx:

Chronic dacryocystitis:

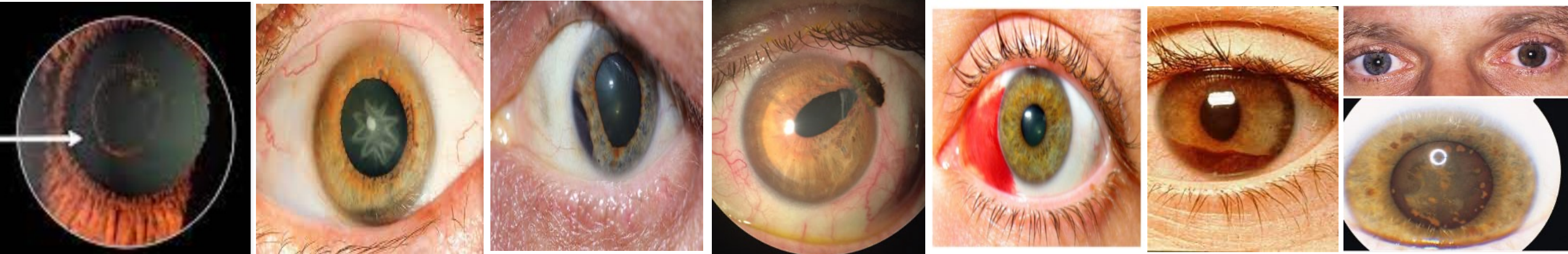


Eyelids



Madarosis:
Ectropion:
Tylosis:

Eye Trauma



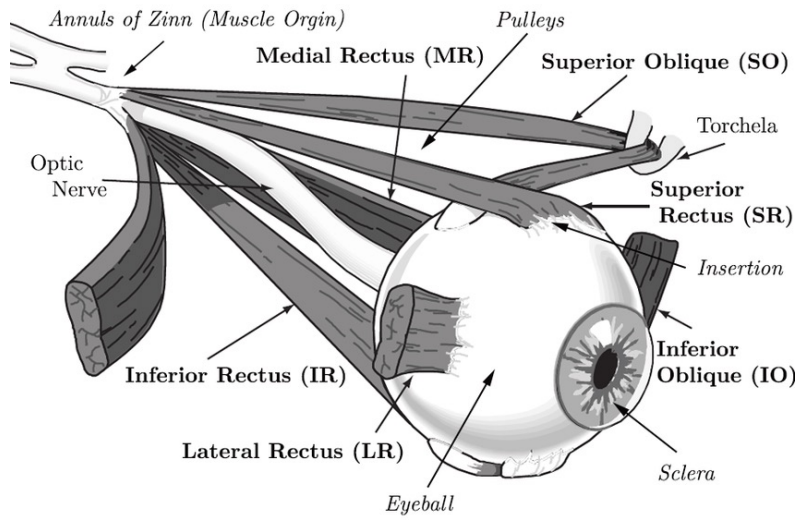
Penetrating trauma
Ciliary body
Granulomatous panuveitis in
C/L eye in 2 weeks
Dalen Fuch nodules

Hudson-Stahli line
Iron deposition around pterygium:
Iron deposition around filtering bleb:

Enucleation
Evisceration
Exenteration

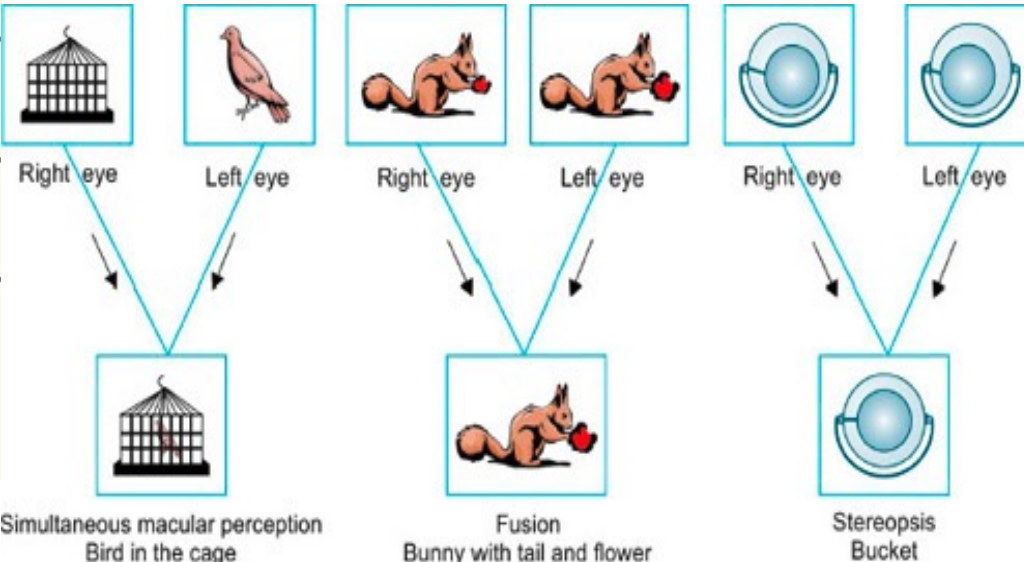
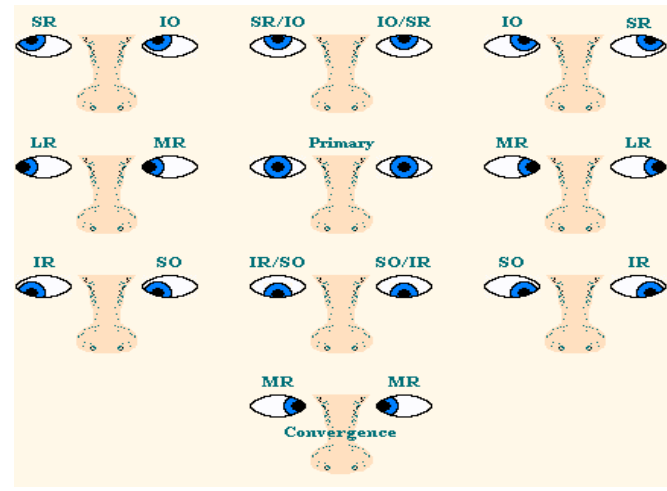
SQUINT – EOM

Misalignment of visual axis
Muscle not arising from Annulus:
Longest muscle:



Muscle	Nerve Supply	Primary Action	Secondary Action	Tertiary Action
MR			-	-
LR			-	-
SR				
IR				
SO				
IO				

HERRING:
SHERRINGTON:



SQUINT – Types

Apparent/Pseudo:

Tropia:

Phoria:

Eso-

Causes:

Exo-

Hyper-

Hypo-

Crossed Diplopia:

Uncrossed Diplopia:

1 = 2 deviation:

2 > 1 deviation:

Forced duction test:



Ptosis + dilated pupil:
Ptosis + constricted pupil:
Ptosis + normal light reflex

Park 3 step test

“IGI and ITO”

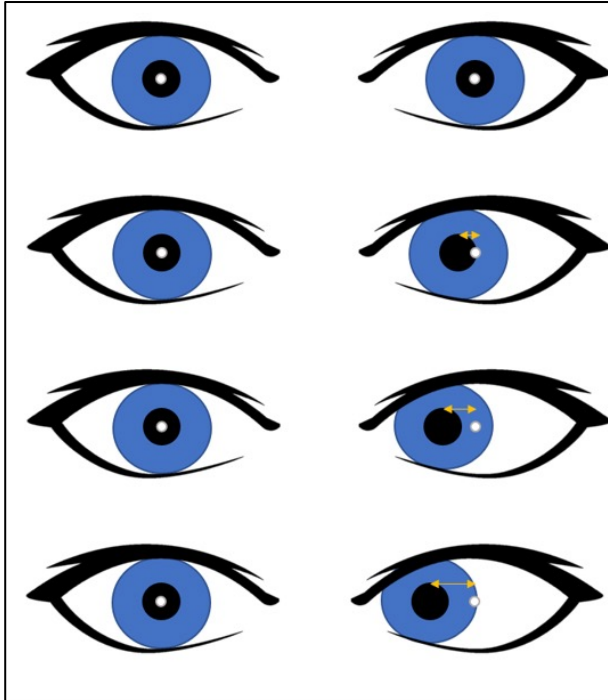
Left hypertropia-Increases on right tilt and right gaze

Right hypotropia-Increases on left tilt and right gaze

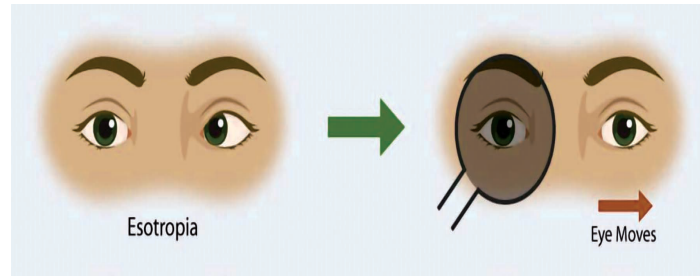
A patient presents with left-sided head tilt which on straightening leads to right hypotropia. This increases on dextroversion and right head tilt. Paralysis of which of the following muscles is involved?

- A) Right superior oblique**
- B) Right superior rectus**
- C) Right inferior oblique**
- D) Left inferior oblique**

SQUINT – TESTS



Hirschberg test:
Amount of squint



Eyes appear normal initially

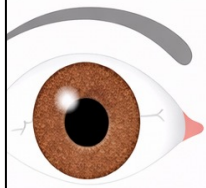

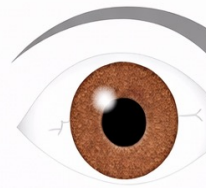




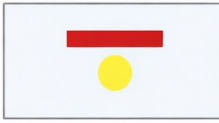
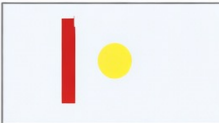



Abnormal eye drifts when covered



Then corrects when uncovered



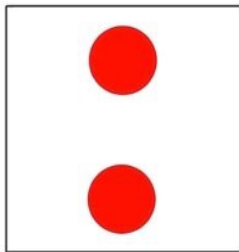
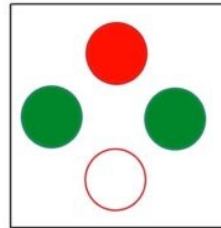
			
Right eye	Left eye	Right eye	Left eye
Orthophoria		Orthophoria	
Esophoria		Hypophoria	
Exophoria		Hyperphoria	

Remove fusion of the two eyes
Lens: Point source -> line perpendicular to axis
Keep Maddox rod // to axis of squint

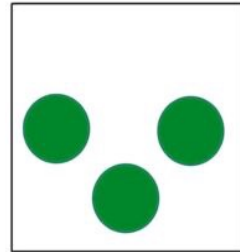
In a patient of left eye exophoria, Maddox rod was placed in front of the right eye, with its axis horizontal. Patient will see bright spot in relation to the red line as:

- A. Horizontal red line with bright spot above it**
- B. Horizontal red line with bright spot below it**
- C. Vertical red line with bright spot right to it**
- D. Vertical red line with bright spot left to it**

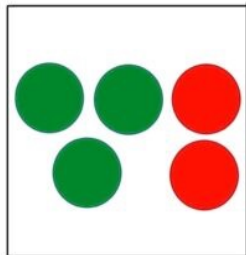
SQUINT – TESTS



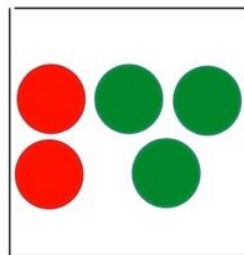
LE SUPPRESSION



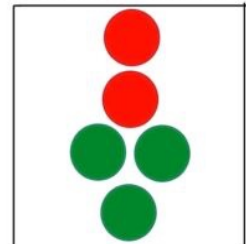
RE SUPPRESSION



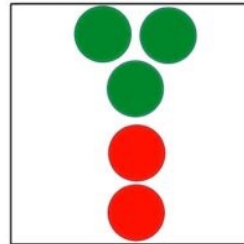
UNCROSSED DIPLOPIA



CROSSED DIPLOPIA



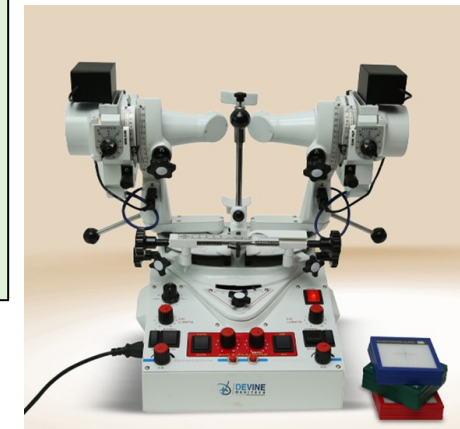
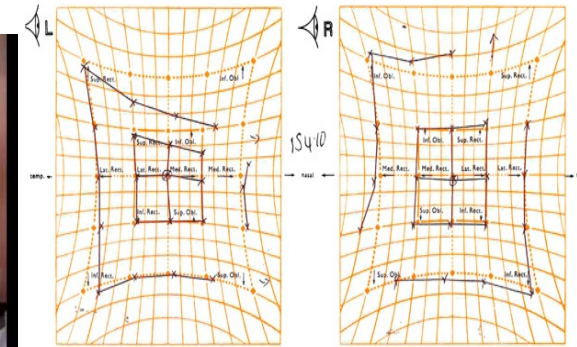
LEFT HYPER DEVIATION



RIGHT HYPER DEVIATION



Most accurate for degree of tropia
 Eso: Base
 Exo: Base
 Hypo: Base
 Hyper: Base



Miscellaneous

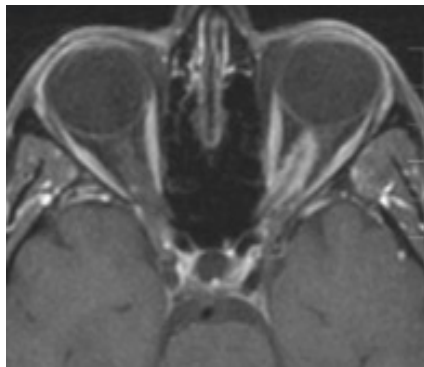


Amblyopia (Lazy eye)
Loss of vision without organic cause
Prevention:
Rx of strabismus: Recession
Resection

**Synkinetic
phenomenon b/w V
and III CN:**

Surgery	Levator Action
Fasanella-Servat	Good (Horner's syndrome) Mild ptosis
Levator Resection	Moderate
Frontalis/ Fascia lata Suspension	Poor (Severe ptosis) Congenital/ Marcus-Gunn ptosis

**Defective elevation in
adduction, normal
elevation in abduction**

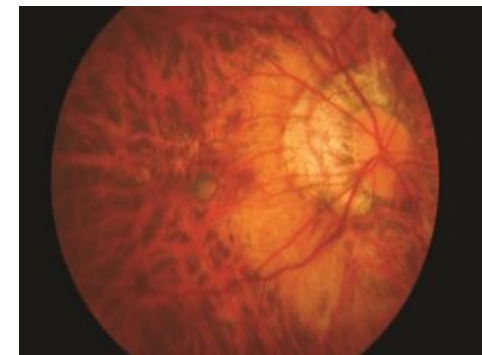
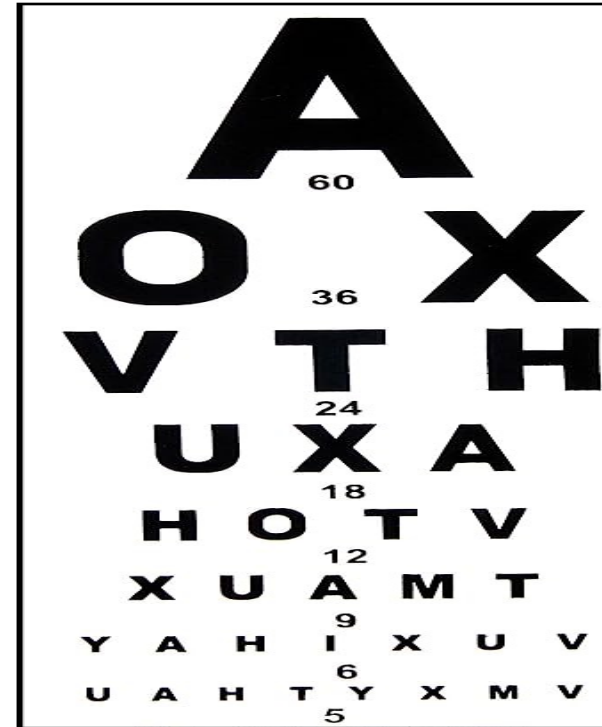


Refractory Errors

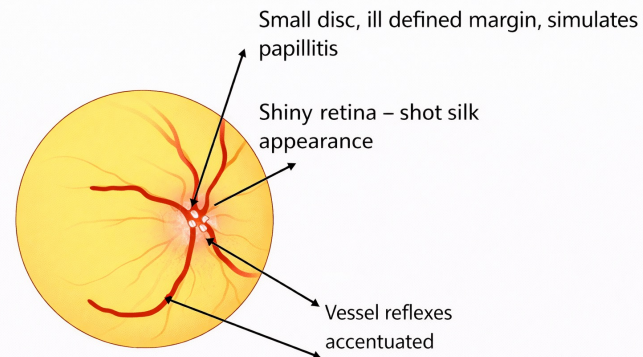
MYOPIA
 Axial length:
 AC:
 Squint:

Axial myopia:
Curvature myopia:
Index myopia:
Positional myopia:

HYPERMETROPIA
 Axial length:
 AC:
 Squint:
 Aphakia



Tigroid appearance
Chorioretinal atrophy
Foster Fuch spots
Lacquer cracks
Rhegmatogenous RD
Post staphyloma



Treatment of myopia :
Radial keratotomy
LASIK: Laser-Assisted In Situ Keratomileusis
CI: <18yrs, unstable power, CT <450um
SMILE: Small Incision Lenticule Extraction
Phakic refractive lens (PRL)/ implantable collamer lens (ICL)



Each letter subtends angle of 5min of arc when viewed from respective distance



Angle subtended by topmost letter when viewed from 6m:

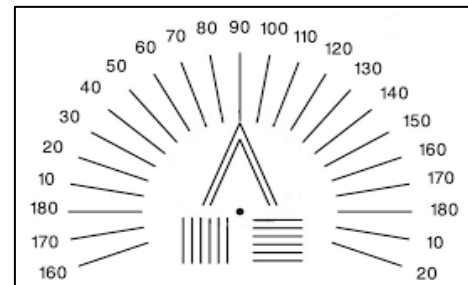
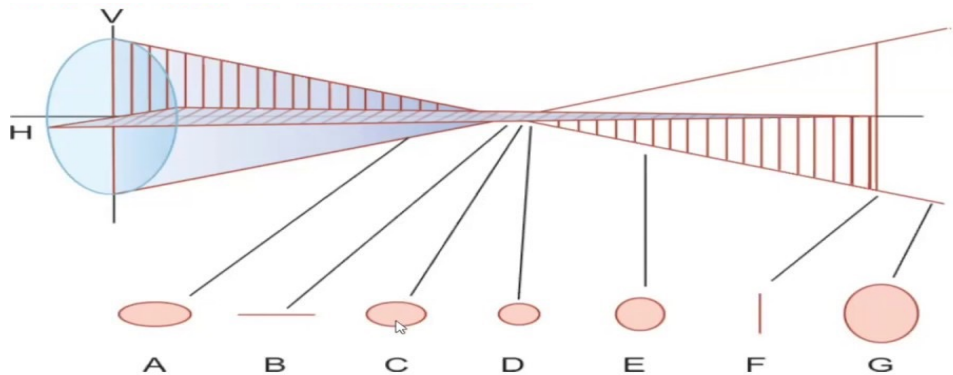
Landolt C chart
ETDRS
Tellers chart, Optokinetic drum

ASTIGMATISM

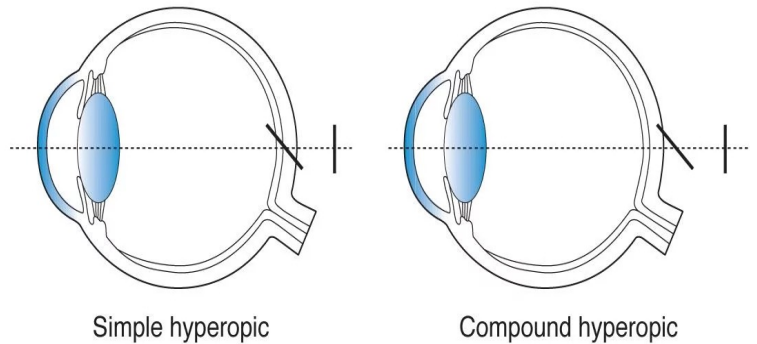
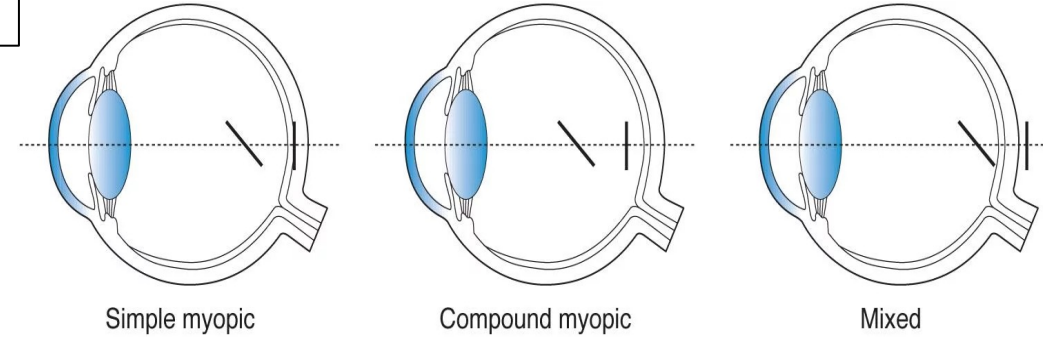
Unequal curvature of eye in different meridians of the eye

Regular: 2 principal meridians perpendicular to each other

Regular astigmatism		180	90
WTR Vertical more curved			
ATR Horizontal more curved			

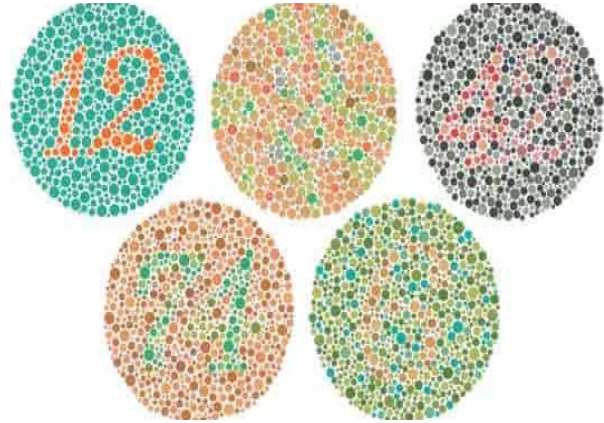


Simple: One meridian emmetropic



-2cyl @180:
-1DS /-2DC @180
-3DS/ -4DC @90:

Miscellaneous



Retinoscopy
Examination distance:
No mvt:
Against:
With:

Prota Deutra Trita
Inheritance:
MC:

Photocoagulation
Double freq Nd-Yag (532nm)
Argon laser

Photoablation
Argon fluoride Excimer
(193nm)

Photodisruption
Nd-Yag (1064nm)

Nd Glass (1053nm)

Clinical presentations-Localisation:
Gradual progressive painless DOV:
Sudden painful red eye + DOV:
Sudden painless DOV:
Floater:
Metamorphopsia:

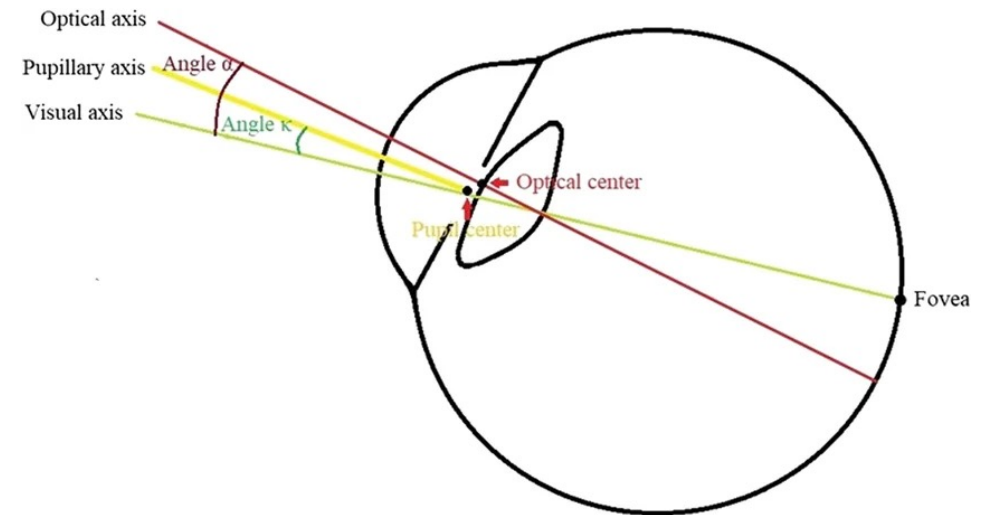
GTCS

Asteroid hyalosis

Synchysis scintillans

Eye drop -> ointment

Chemical injury to eye:



$\geq 6/18$
 $< 6/18 - 6/60$
 $< 6/60 - 3/60$
 $< 3/60 - 1/60$
 $< 1/60 - LP+$
NLP

Alpha – between visual axis and optical axis
Gamma – between fixation axis and optical axis
Kappa – between visual axis and pupillary plane

