

## Cataract (Definition)

*Opacification of the lens of the eye

* It is the most common cause of treatable blindness in the world


Lens clouded by cataract


## Age Related Cataract

$\square$ The main types of age-related cataracts are :

1. Nuclear sclerosis
2. Cortical
3. Posterior subcapsular.




## 3) Posterior subcapsular

$\square$ Forms on the back of the lens, on the inner surface of the posterior capsule bag.
$\square$ These cataracts tend to occur in patients on steroids, with diabetes, and those with history of ocular inflammation.
$\square$ The opacity looks like breadcrumbs or sand sprinkled onto the back of the lens. This posterior location creates significant vision difficulty despite appearing innocuous on slit-lamp exam.


Phacoemulsification

1. Capsulorhexis
2. Sculpting of nucleus
3. Emulsification of each quadrant

4. Hydrodissection
5. Cracking of nucleus
6. Cortical cleanup and insertion of IOL

## Extracapsular cataract extraction

1. Anterior capsulotomy
2. Expression of nucleus
3. Care not to aspirate posterior capsule accidentally

4. Completion of incision
5. Cortical cleanup
6. Polishing of posterior capsule, if appropriate

## Extracapsular cataract extraction ( cont. )

7. Injection of viscoelastic substance
8. Insertion of inferior haptic and optic
9. Placement of haptics into capsular bag and not into ciliary sulcus

10. Grasping of IOL and coating with viscoelastic substance
11. Insertion of superior haptic
12. Dialing of IOL into horizontal position

## Aqueous outflow



With the loss of nerve fibers from glaucoma, the cup becomes progressively larger because there is less space occupied by the remaining nerve fibers

The optic nerve is divided into tenths and the cup is compared to the entire optic nerve (optic disc) to obtain the cup-to-disc ratio.

The normal c/d ratio is 0.3(The C/D ratio here is 0.4)



## Progression of glaucomatous cupping


a. Normal (c:d ratio 0.2)
b. Concentric enlargement (c:d ratio 0.5)
c. Inferior expansion with retinal nerve fibre loss
d. Superior expansion with retinal nerve fibre loss
e. Advanced cupping with nasal displacement of vessels
f. Total cupping with loss of all retinal nerve fibres

## Primary Glaucoma

- Classification of the primary glaucomas is based on whether or not the iris is:
- Clear of the trabecular meshwork (open angle) $\rightarrow$ (POAG )
- Covering the meshwork-(closed angle) $\rightarrow$ (PACG )


ANGLE: the iridocorneal angle

OPEN AND CLOSED ANGLE GLAUCOMA


## How to examine the ANGLE

- Using special lenses which allows visualization of the angle structure to determine whether the angle configuration is closed or open Goniolenses.



## Technique




The appearance of a trabeculectomy bleb.


ACUTE POSTERIOR VITREOUS DETACHMENT: (A) SYNCHISIS; (B) UNCOMPLICATED POSTERIOR VITREOUS DETACHMENT; (C) RETINALTHR FORMATION AND VITREOUS HAEMORRHAGE; (D) AVULSION OF A RETR (fふ̧L BLOOD VESSEL AND VITREOUS HAEMORRHAGE

- A doctor can usually diagnose macular degeneration by examining the eyes with an ophthalmoscope or slit lamp. Sometimes fluorescein angiography is used to determine the diagnosis

dry AMD, note the discrete scattered yellowish sub-retinal drusen

wet AMD, note the small haemorrhage associated with the subretinal membrane


## Static traction:

-It plays an important role in the pathogenesis of tractional RD and proliferative vitreoretinopathy.
-It can be: Tangential, Anteroposterior and Bridging traction.



## DYNAMIC VITREORETINAL TRACTIONS:



Normal Vitreoretinal adhesions

TRACTIONAL RD IN PROLIFERATIVE DIABETIC RETINOPATHY



Tractional RD in proliferative diabetic retinopathy


INFERIOR TRACTIONAL RETINAL DETACHMENT


RETINAL TEAR WITH RETINAL DETACHMENT


## EXUDATIVE RETINAL DETACHMENT IN VKH



## CHOROIDAL DETACHMENT




RETINOSCHISIS



Pupil


## Vascular Tunic -- Muscles of the Iris



Bright light


Normal light

Radial muscles


Dim light

- Constrictor pupillae (circular) are innervated by parasympathetic fibers while Dilator pupillae (radial) are innervated by sympathetic fibers.
- Response varies with different levels of light




Normal<br>Response to Light



## Positive RAPD of Right Eye




Sphincter
pupillae m


Ciliary ganglion

Pretectal nucleus

Superior cervical ganglion

## Parasympathetic Pathway

 Sympathetic PathwayMiosis
Ptosis
Apparent enophthalmos

## Horner's

- Characterized by the classic triad of
- Miosis (constricted pupil)
- Partial ptosis
- Loss of hemifacial sweating
 ( anhidrosis).


Laxmi
eye institute

## Brain stem-ventral

 view

4 CN are in above pons (I, II, III, IV).
4 CN are in pons (V, VI, VII, VIII).
4 CN are in medulla (IX, X, XI, XII).
4 CN nuclei are medial (III, IV, VI, XII). "Factors of 12 , except 1 and 2 ."

Supraorbital notch

Superior orbital fissure


Inferior orbital fissure

Whitnall's tubercle

Infraorbital foramen


## Ocular motility



CNVI innervates the Lateral Rectus. CN IV innervates the Superior Oblique.
CN III innervates the Rest.
The "chemical formula" $\mathrm{LR}_{6} \mathrm{SO}_{4} \mathrm{R}_{3}$.
The superior oblique abducts, intorts, and depresses while adducted.


Obliques go Opposite (leff SO and IO tested with patient looking right).
IOU: 10 tested looking Up.

## CN III. Oculomotor nerve

Origin/Course


Photograph of the patient's extraocular movements on forward gaze (A), right gaze (B), and left gaze (C)

## ISOLATED NERVEPALSIES



Right gaze


Upgaze


## CN IV. Trochlear nerve

 Origin/Course

## FOURTH CRANIAL NERVE

 PALSIESWeakness of the muscle innervated by the 4th (trochlear) nerve

- These palsies are often difficult
 to detect because they produce the eye movement abnormality.
- They affect predominantly when the eve is turned inward (Intorsion).



## Originates in the Abducens Nucleus $\rightarrow$

"PontoMedullary Junction"
Continues anteriorly through the Cavernous sinus
Then continues to the lateral portion of the "Superior Orbital Fissure"


Bilateral fth nerve palsy:

"Look to your RIGHT

Look to your LEFT"


Fig. 19.71 Acute left sixth nerve palsy in a child. (A) Left esotropia in the primary position; (B) marked limitation of left abduction


The normal optic nerve head has distinct margins, a pinkish rim and usually a white central cup "CDR=0.3"


## Optic Disc Swelling

a) A normal disc
(B) A SWOLLEN DISC SECONDARY TO RAISED INTRACRANIAL PRESSURE. NOTE THE LACK OF A SHARP OUTLINE TO THE DISC AND THE DILATED CAPILLARIES ON THE DISC.

(c) The appearance of optic disc drüsen; note how the solid yellow lesions cause irregularity of the disc margin

(d) Myelination of the nerve fibers around the nerve head may be mistaken for a swollen optic disc

(e) A myopic optic disc. Note the extensive peripay may atrophy.



## Pale with swollen disk:

suggested GCA particularly if associated with visual loss

Pale,
featureless disc
$=$ optic atrophy
(ischaemia, MS etc)

(a)

(b)

## Normal Optic Disc



Optic Neuritis


Neuroretinitis

## Advanced techniques Slit Lamp





## Optical Coherence Tomography: OCT

Scan angle: $0^{\circ} \quad 5$ pacing: 0.25 mm Length: 5 mm


## Ultrasound: B-scan

## Ultrasound: A-scan



## Fluorescein Angiography



Arterial phase


Venous phase


Early venous phase


Late phase


Fig. 4 Comeal contour chart produced by computed tomography.


Fig. 2 An Ishihara pseudoisochromatic plate.

## MINOR TRAUMA:

- Conjunctival FB:
- SUBTARSAL FOREIGN BODY: No ocular examination is complete until the upper eyelid is everted and closely inspected.
- Linear epithelial defects is suggestive of a foreign body under the eye lid

- Corneal FB:
- Corneal foreign bodies and rust rings are best removed with a sterile disposable needle (19 to 23 G ).
- Do not attempt central or deep foreign bodies as a slit lamp is required to avoid excessive trauma.
- Instil antibiotic ointment and pad for 24 hours.
- Review daily until healed, or referred to ophthalmologist.


-Damage from alkali injuries is usually worse and carries a poor prognosis(Fig. 2).


## ALKALI BURN UNTREATED




## SUBCONJUNCTIVAL HEMORRHAGE

- Can occur secondary to blunt trauma or can be spontaneous
- Usually benign and self-limited.
- No treatment is required, Lubrication if foreign body sensation
- But be aware of rupture globe.



## SUBCONJUNCTIVAL HEMORRHAGE



## SUBCONJUNCTIVAL HEMORRHAGE





- RETINAL DETACHMENT. BEWARE OF FLOATERS, FLASHES AND FIELD DEFECTS. URGENT REFERRAL IS MANDATORY.


CHOROIDAL RUPTURE


Commotio Retinae w/ retinal hgs

## RUPTURE OF GLOBE



## OPEN GLOBE

- Perforating corneal laceration



## PENETRATING TRAUMA

- Do not remove penetrating foreign bodies
- Minimize additional damage (shield)
- NPO
- Tetanus
- Antibiotics
- REFER

- Penetrating injury with nail gun


- Severe trauma that resulted in a scleral rupture with delivery of the lens

- Cataractous lens following penetrating trauma

INTRAOCULAR FOREIGN BODY


## INTRAOCULAR FOREIGN BODY

- CT localizes best
- Antibiotics (IV or PO)
- Refer



## OPEN GLOBE-TREATMENT

- When suspect an open globe, one shall protect the eye from further injury by covering the eye with a shield or any device that can protect the eye.
- Minimize additional damage
- shield (not patch)
- avoid valsalva
- Do not instill any medication to the eye.
- Do not attempt to remove anything from the wound, and consult the opthalmologist.
- "Prepare for repair"
- NPO



## LID LACERATIONS



## LID LACERATION

- REFER for location
- medial
- margin



## LID LACERATION

- REFER for
- Depth
- Extensive tissue loss



## ORBITAL BLOW OUT FRACTURE

- Abnormal motility -> DOUBLE VISION
- May need repair (if persistent)



## ORBITAL BLOW OUT FRACTURE

- Crepitus
- Paresthesia (cheek, gum)



## Bacterial keratitis



## Herpes simplex keratitis



## Herpes Zoster Keratitis

Late changes: severe dry eye with
macrodendrite
s , post herpetic neuralgia
Treatment: oral Acyclovir of proven benefit to reduce complications, topical antivirals not indicated, lubrication, treat
complications


## Episcleritis



## Episcleritis



## Episcleritis



- Treatment
- Mild cases may need no specific therapy but if discomfort is annoying, topical steroids and/or topical non-steroidal anti-inflammatory drugs (NSAIDs) may be helpful.
- Unresponsive recurrent cases, which are rare, require systemic flurbiprofen ( 100 mg three times daily), taken at the first symptom of recurrence in order to abort an attack


## ANTERIOR NON-NECROTIZING SCLERITIS

- Diffuse scleritis is characterized by zoidespread - Nodular scleritis



## ANTERIOR NON-NECROTIZING SCLERITIS

- Nodular scleritis



## Anterior Necrotizing scleritis with inflammation

- Distortion or occlusion of blood vessels
- Development of scleral necrosis



## Anterior Necrotizing scleritis with inflammation

- Development of scleral necrosis
- Distortion or occlusion of blood vessels



Anterior Necrotizing scleritis without inflammation

- scleromalacia perforans



## POSTERIOR SCLERITIS

- External signs may include eyelid oedema, proptosis anc ophthalmoplegia.

- Ophthalmoscopy shows Disc swelling, macular oedema and exudative retinal detachment .
- Ultrasonography shows thickening of the posterior sclera with fluid in Tenon space
- CT will also demonstrate posterior scleral thickening .
- Treatment
- Elderly patients with associated systemic disease are treated in the same way as those with necrotizing anterior scleritis.
- Young patients without associated systemic disease usually respond well to NSAIDs.


## UVEITIS

## Ciliary injection:



## Keratic precipitates:



## UVEITIS

## Iris Nodules



UVEITIS

Hypopyon


## UVEITIS

## Flare:



UVEITIS

## Posterior synechia




## Bacterial

## Conjunctivitis

- Common pathogens:
- Staphylococcus coagulase or noncoagulase positive,
- Streptococcus
- Hemophilus influenzae,
- Pseudomonas sp.



## Hyperpurulent Conjunctivitis

- caused by gonococcus
- URGENT REFERRAL for intensive IV \& topical treatment



## Viral Conjunctivitis





## CHLAMYDIAL CONJUNCTIVITIS

Signs include: edematous conjunctiva, mucopurulent discharge, papilla then follicular reaction, non-tender lymphadenopathy.
: Topical therapy is with tetracycline ointment four times daily for 6 weeks.

- Systemic therapy can be with one of the following:
- Doxycycline either 300 mg weekly for 3 weeks or 100 mg daily for l-2 weeks.
- Tetracycline 250 mg four times daily for 6 weeks.
- Erythromycin 250 mg four times daily for 6 weeks if tetracycline is inappropriate.

Neonatal chlamydial conjunctivitis

- Treatment is with topical tetracycline and oral erythromycin $25 \mathrm{mg} / \mathrm{kg}$ body weight twice daily for 14 days.


## Allergic Conjunctivitis




- Pingueculum
- Treatment: lubricant and/or vasoconstrictor
- Pterygium



## - Subconjunctival Haemorrhage :



- Lensectomy + posterior capsulotomy + anterior vitrectomy.


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## The WAGR

 Syndrome

## Foveal Hypoplasia

- Causes: Aniridia and Albinism


- Pseudostrabismus




## Hirschberg's test

Amount of deviation: note location of corneal light reflex

$$
1 \mathrm{~mm}=7^{\circ} \text { or } 15 \Delta
$$



Reflex at border of pupil $=15^{\circ}$


Reflex at limbus $=45^{\circ}$

## Hirschberg light Test

- Used to test tropia but not phoria.
- It is performed by shining a light in the person's eyes and observing the light reflection from the corneas.
- Nasal corneal light reflex: exotropia
- Temporal corneal light reflex: esotropia
- Inferior corneal light reflex: hypertropia
- Superior corneal light reflex: hypotropia



NON-ISCHAEMIC CENTRAL RETINAL VEIN OCCLUSION


FLUORESCEIN ANGIOGRAM OF NON-ISCHAEMIC CENTRAL RETINAL VEIN OCCLUSION SHOWING CYSTOID MACULAR OEDEMA BUT GOOD PERFUSION

## ISCHEMIC CRVO:

- Afferent papillary defect, more hemorrhages and hard exudates(Stormy appearance) with more severe optic disc swelling


FLUORESCEIN ANGIOGRAM OF LSCHAEMIC CENTRAL RETINAL VEIN OCCLUSION SHOWING EXTENSIVE AREAS OF CAPILLARY NON-PERFUSION

## MANAGEMENT

- Macular edema:
- Anti-VEGF
- Steroids
- Argon laser
- Vitrectomy in resistant cases
- Neovascularizations:
- Panretinal Photocoagulation
- Vitrectomy for advanced stage
- Specific glaucoma treatment for NVG

