

## 4. Diseases of the Sclera

### Q Give an account on the scleral foramina

**Answer**

#### **Scleral foramina:**

- (A) Large foramina (2 in number)
- (B) Small multiple foramina

#### **(A) Large foramina (two in number):**

##### **1. Posterior scleral foramen.**

- Lies 3 mm medial and 1 mm above the posterior pole of the eye.
- It is the site of exit of optic nerve.
- The optic nerve fibers pierce the sclera which is weakened and has a sieve-like appearance and is known as Lamina cribrosa.
- The sclera is fused at the edge of the posterior scleral foramen with the dural and arachnoid sheaths of optic nerve.
- The central opening of lamina cribrosa is large and transmit the central retinal vessels.

##### **2. Anterior scleral foramen:**

The cornea fits in the anterior scleral foramen like a watch glass.

#### **(B) Multiple small foramina:**

##### **► Three groups exist:**

##### **a) Anterior:**

Located at the insertion of recti muscles or immediately in front of it. Transmit branches of the anterior ciliary arteries (7 in number: 2 from each rectus muscle except the L.R. which only has one).

##### **b) Middle:**

About 4 mm behind the equator and number 4 or 5. These are for the exit of vortex veins, which drains the uveal tract, mainly choroid.

##### **c) Posterior:**

Are small, numerous and located around the optic nerve. They transmit the long and the short posterior ciliary nerves and vessels.

### Q Differentiate between episcleritis and phlyctenular conjunctivitis

#### **D.D.:**

Phlyctenular kerato conjunctivitis (**Table** ).

[Table](#) : DD of nodular episcleritis and phlycten

Character:	Nodular episcleritis	Phlycten
<b>(1) Incidence:</b> 1) Age: 2) Sex:	Old age More in females	Children. Equal.
<b>(2) History:</b> <b>(3) Signs:</b> 1) Site.	Long.  Some distance from the limbus.	Short.  At the limbus.

2) Colour:	Purple (deep vessels).	Grey.
3) Level:	Subconjunctival (deep).	Conjunctival (superficial).
4) Movement:	Fixed to sclera and conjunctiva moves over it.	Moves with conjunctiva.
5) Tenderness:	Very tender.	Not tender.
6) Ulceration:	Never occurs.	Occurs.
7) Suppuration:	Never occurs.	Occurs.
8) Adrenaline test:	No blanching (deep vessels).	Blanching (superficial vessels)
<b>(4) Complications:</b>	Rare (scleritis).	Common (corneal).
<b>(5) Prognosis:</b>	May be serious (cause).	Good.

**Q Mention the classification, clinical picture, and complications of scleritis**

**Answer**

**Classification (Clinical Types) of Scleritis:**

**(1) Anterior Scleritis**

**a) Non-necrotizing scleritis:**

- i. Nodular
- ii. Diffuse

**b) Necrotizing scleritis (with & without pain) :**

**(2) Posterior scleritis**

**(I) Anterior Scleritis**

**(A) Anterior Non-necrotizing Scleritis:**

• **Clinical Picture:**

**1. Symptoms: Pain** is common symptom and may radiate to the frontal or maxillary regions, **marked tenderness**.

**2. Signs:**

**a) Nodular Scleritis: A localized nodular lesion with the following characters:**

- **Less circumscribed** than episcleritis.
- **Dark red or violet** at first, later it becomes pale purple.
- It may surround the cornea. (**Annular scleritis**).
- **Never ulcerate**, the cornea and iris may be involved.

**b) Diffuse Scleritis:**

- **Widespread inflammation**

- **No ulceration**, but much absorption of the sclera causing marked **thinning** and **staphyloma**.

• **Treatment:**

- **Non-steroidal Anti-inflammatory drugs (NSAID)** e.g. indomethacin 100 mg daily for 4 days then 75 mg daily until resolution.

## **(B) Anterior Necrotizing Scleritis (with & without Pain):**

### **(i) Anterior Necrotizing Scleritis with pain:**

- **Clinical Features:**

- a. **Presentation (Symptoms):** with **severe pain**.

- b. **Signs:** Localized patches of scleral necrosis exposing the uvea..

- **Treatment:**

- Systemic steroids or

- Immunosuppressive drugs e.g. cyclophosphamide.

### **(ii) Anterior Necrotizing Scleritis without Pain (Scleromalacia Perforans):**

- **Clinical Features:**

- c. **Presentation (Symptoms):** usually in females with long standing generalized seropositive **rheumatoid arthritis** with an asymptomatic (**painless**) **dark scleral patch** due to scleral atrophy.

- d. **Signs:** Large patches of scleral necrosis exposing the uvea..

- **Treatment:**

- 1. **Medical treatment:** usually not effective.

- 2. **Surgical treatment:**

- 1- **Extreme thinning of the cornea and sclera:**

- Corneal grafting

- Scleral grafting covered by conjunctiva.

- Fascia lata or periosteum are more resistant to the melting process.

- 2- **Extreme corneal marginal ulceration or keratolysis:** may require corneal grafting (usually lamellar patch graft).

- **Complications of anterior scleritis:**

- 1- **Uveitis** with its complications, especially cyclitis and anterior choroiditis.

- 2- **Sclerosing keratitis.** (triangular with rounded apex, with little or no corneal vascularization, ulceration never occur, partial clearing occur, pupillary area usually escape), the densest parts usually persists as a bluish clouds.

- 3- **Secondary glaucoma.**

- 4- **Ectasia** of the thinned sclera (ciliary staphyloma).

- 5. **Loss of vision.**

- 6- **Brawny scleritis:** diffuse form with involvement of extra ocular muscles, with marked oedema and diffuse redness, chronic course. Loss of the eye, may occur.

- 7- The most serious complication is **keratolysis** wherein the stroma melts away with Descematocele formation.

- 8- Corneal and linnbal **guttering** with thinning and ectasia.

- 9- **Complicated Cataract.**

- 10. **Retinal detachment.**

- 11 - **Papilloedema.**

## **(II) Posterior Scleritis**

It is actually scleral and Tenon's capsule inflammation behind the equator.  
**leading to:**

- 1- Moderate pain and oedema of lids.**
- 2- Proptosis.**
- 3- Limitation of ocular movement with diplopia.**

• **Complications of posterior scleritis:**

1. Choroidal effusion
2. Exudative RD
3. Macular edema
4. Optic disc edema (swelling)

**Q What are the types of ocular staphylomata**

**Answer**

• **Definition:**

A staphyloma is a bulge of the outer coat of the eye, lined by atrophic uveal tissue.

• **Clinical Types:**

It may be **corneal or scleral**

**A. Corneal Staphyloma:**

- It is a bulge of a corneal scar with incarcerated atrophic iris tissue.
- It is caused by perforated corneal ulcer or wound with iris prolapse
- The surface appears lobulated bluish in colour.
- On healing lead to glaucoma if a sufficient area of the angle is blocked by peripheral anterior synechiae.
- If the condition is neglected blindness follow.

- **Types:** it may be **partial or total.**

**1) Partial corneal staphyloma :** Conical bulging of a big corneal scar with iris in carceration due to large corneal perforation with iris prolapse.

**2) Total corneal Staphyloma :** Hemispherical bulging of the pseudo-cornea due to sloughing of the whole cornea → sloughing of the whole cornea.

**B. Scleral staphylomata:**

• ***These may be:***

**1. Anterior scleral staphyloma**

- where the sclera is weakened by the ciliary B.vs. and schlemm's canal.
- ***This includes:***

**a) Intercalary staphyloma:**

- Occurs within 2-3 mm zone concentric with the limbus.
- This part is weakened by schlemm's canal.

- The staphyloma is **lined by atrophic iris root and peripheral anterior synechiae**.

- **It is in front of the perforating branches of the anterior ciliary arteries.**

- **Causes:** It is caused usually by prolonged rise of IOP in the degenerative stage of the absolute **glaucoma**

**b) Ciliary staphyloma:**

- It is a bulge of the sclera, situated behind the limbus, **lined by atrophic ciliary body**.

- It is **behind the perforating branches of the anterior ciliary vessels**.

- Its surface is irregular, bluish in colour.

- It may surround the cornea (**Ring staphyloma**).

- It is slowly progressive and may rupture.

- **Causes:** it is caused by weakness of the sclera following **scleritis or injury**, usually associated with some rise of the IOP, and also caused by prolonged rise of I.O.P. in the degenerative stage of **absolute glaucoma**.

**2. Equatorial staphyloma:**

- Occurs at that part of the eyeball which is weakened by the passage of the vortex veins **usually between recti muscles, lined by atrophic chorioretinal tissue**. (usually seen after excision of the globe).

- **Causes:** Caused by **absolute glaucoma**.

**3. Posterior Staphyloma:**

- Which occurs at the posterior pole of the eye, usually on the **temporal** side of the optic disc, or may surround the optic disc (**peripapillary staphyloma**).

- **Cause:** Caused primarily by the scleral degeneration of **high degree of axial myopia**.

- **Diagnosis:** Can be diagnosed **ophthalmoscopically** (the B.v.s. being at two levels). **The best diagnostic method is by ultrasonography**.

• **Treatment of Ocular Staphylomata** : Depends upon the cause.

(1) **Treatment of the cause:** If treatable.

(2) **Treatment of the staphyloma:**

1) Partial corneal staphyloma: Penetrating keratoplasty.

2) Scleral staphyloma: Scleral graft especially small equatorial staphyloma;

(3) **Treatment of secondary glaucoma:** Filtering operation.

(4) **Treatment of a blind painful eye:** Enucleation.

**Q What are the causes of blue sclera**

**Answer**

**BLUE SCLERA**

• **Definition**

Abnormal discolouration of thin sclera due to uveal tissue showing through it (as sclera is thin).

• **Aetiology:**

(1) **Osteogenesis imperfecta**: **A hereditary syndrome characterized by:**

1) Fragile bones.

2) Deafness (due to otosclerosis).

3) Blue sclera (due to thin sclera).

**(2) Thin sclera:**

- 1) In children.
- 2) In high myopia.
- 3) In scleral staphyloma.
- 4) In buphthalmos.

