

OCULAR DISEASE

SCLERITIS



Introduction

- Scleritis refers to a chronic inflammation of the sclera proper.
- It is a comparatively serious disease which may cause visual impairment and even loss of the eye if treated inadequately.
- It usually occurs in elderly patients (40-70 years) involving females more than the males.

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Etiology

Autoimmune collagen disorders

- 1. Rheumatoid arthritis.
- 2. about 5% cases of scleritis are associated with connective tissue disease.
- 3. About 0.5 percent of patients (1 in 200) suffering from seropositive rheumatoid arthritis develop scleritis.
- 4. Other associated collagen disorders are
- > Wegener's granulomatosis,
- > polyarteritis nodosa (PAN),
- Systemic lupus erythematosus (SLE)



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- Metabolic disorders like gout and thyrotoxicosis have also been reported to be associated with scleritis.
- 3. Some infections, particularly herpes zoster ophthalmicus, chronic staphylococcal and streptococcal infection.
- 4. *Granulomatous diseases like tuberculosis,* syphilis, sarcoidosis, leprosy can also cause scleritis.
- 5. *Miscellaneous conditions like irradiation, chemical* burns, Vogt-Koyanagi-Harada syndrome, Behcet's disease



Pathology

Histopathological changes are that of a chronic granulomatous disorder characterised by fibrinoid necrosis, destruction of collagen together with infiltration by polymorphonuclear cells, lymphocytes, plasma cells and macrophages



Classification

- I. Anterior scleritis (98%) 1. Non-necrotizing scleritis (85%) (a) Diffuse (b) Nodular 2. Necrotizing scleritis (13%) (a) with inflammation (b) without inflammation (scleromalacia
 - perforans)
- II. Posterior scleritis (2%)



Clinical features-Symptoms.

- Patients complain of moderate to severe
- pain which is deep and boring in character and often wakes the patient early in the morning.
- Ocular pain radiates to the jaw and temple.
- Localised or diffuse redness,
- Mild to severe photophobia and lacrimation.
- Occasionally there occurs diminution of vision.



Signs

1. Non-necrotizing anterior diffuse scleritis: Widespread inflammation involving a quadrant or more of the anterior sclera. The involved area is raised and salmon pink to purple in colour





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2. Non-necrotizing anterior nodular scleritis. It is characterised by one or two hard, purplish elevated scleral nodules, usually situated near the

limbus



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3. Anterior necrotizing scleritis with inflammation.

- localised inflammation associated with areas of infarction due to vasculitis.
- The affected necrosed area is thinned out and sclera becomes transparent and ectatic with uveal tissue shining through it. It is usually associated with anterior

uveitis.





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4. Anterior necrotizing scleritis without inflammation (scleromalacia perforans). This specific entity typically occurs in elderly females usually suffering from long-standing rheumatoid arthritis.

 It is characterised by development of yellowish patch of melting sclera





Complications

 These are quite common with necrotizing scleritis and include
 Sclerosing keratitis
 keratolysis

- Complicated cataract
- Secondary glaucoma.



Investigations 1. TLC, DLC and ESR

- Serum levels of complement (C3), immune complexes, rheumatoid factor, antinuclear Antibodies
- 3. FTA ABS, VDRL for syphilis.
- 4. Serum uric acid for gout.
- 5. Urine analysis.
- 6. Mantoux test.
- 7. X-rays of chest, paranasal sinuses, sacroiliac joint and orbit to rule out foreign body especially in patients with nodular scleritis.



Treatment

- (A) Non-necrotising scleritis. It is treated by topical steroid eyedrops and systemic indomethacin 100 mg daily for a day and then 75 mg daily until inflammation resolves.
- (B) Necrotising scleritis. It is treated by topical steroids and heavy doses of oral steroids tapered slowly.
- In non-responsive cases, immuno-suppressive agents like methotrexate or cyclophosphamide may be required.
- Subconjunctival steroids are contraindicated because they may lead to scleral thinning and perforation.