

NOTES **EYE INFECTIONS**

GENERALLY, WHAT ARE THEY?

PATHOLOGY & CAUSES

 Ocular disorders with infectious, noninfectious etiologies → inflammation, damage to eye structures

RISK FACTORS

 Immunocompromised state, contact with infectious agent, ocular trauma, certain systemic diseases

COMPLICATIONS

• Range from benign, self-limiting to visionthreatening infections

SIGNS & SYMPTOMS

Structural damage, functional impairment

DIAGNOSIS

DIAGNOSTIC IMAGING

Fundoscopy

CT scan/MRI

• Orbits, sinuses

LAB RESULTS

• Giemsa/Gram stains; cultures

OTHER DIAGNOSTICS

Snellen chart

TREATMENT

MEDICATIONS

Antimicrobials

OTHER INTERVENTIONS

Address comorbidities

CHALAZION

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PATHOLOGY & CAUSES

- Firm, painless lipogranulomatous inflammatory lump in eyelid; caused by blockage of ocular sebaceous glands
 - Deep chalazion: inflammation of meibomian sebaceous glands
 - Superficial chalazion: inflammation of Zeis sebaceous glands
- Gland obstruction → impissation (decreased flow of secretions) → granulomatous inflammatory response \rightarrow lipogranuloma inflammation → lesion forms on upper (most common)/lower eyelid
- Slow growing; may persist for weeks/ months; deeper within eyelid than hordeolum (stye)

RISK FACTORS

• Rosacea, seborrhea, blepharitis, inflamed hordeolum

COMPLICATIONS

- If large chalazion presses on cornea \rightarrow visual changes
- Recurring chalazion: may signal carcinoma (rare)

SIGNS & SYMPTOMS

• Eyelid erythema; swelling; firm, nodular, rubbery consistency

DIAGNOSIS

OTHER DIAGNOSTICS

- Clinical history, physical examination
- Histological examination: chalazia may indicate eyelid carcinoma

Determine status of meibomian glands;

may demonstrate diffuse inspissation of yellowish contents from eyelid margin orifices



Figure 76.1 A chalazion of the left upper eyelid.

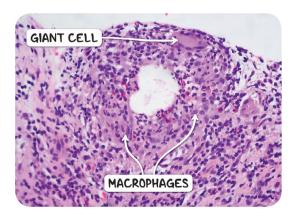


Figure 76.2 The histological appearance of a chalazion. There is granulomatous inflammation with giant cells, numerous macrophages as well as neutrophils and eosinophils surrounding a nidus of lipid.

TREATMENT

MEDICATIONS

 Recalcitrant chalazia: intralesional steroid injection

SURGERY

• Recalcitrant chalazia: incision, curettage

OTHER INTERVENTIONS

- Warm, wet compresses encourage drainage
- Ocular cleansing pads applied to eyelid margin
- Treat comorbidities (e.g. blepharitis, rosacea)
- Small chalazion may resolve on own

CHORIORETINITIS

osms.it/chorioretinitis

PATHOLOGY & CAUSES

 Inflammation of choroid, retina; AKA posterior uveitis

CAUSES

Infectious

- Bacterial: tuberculosis, syphilis
- Viral: cytomegalovirus, West Nile virus, herpes simplex virus (HSV) 1
- Parasitic: toxoplasmosis, onchocerciasis
- Fungal: Candida albicans

Noninfectious

 Sarcoidosis, Behçet's disease, traumatic chorioretinitis

RISK FACTORS

 Immunodeficiency, contact with infectious agent, traumatic eye injury, systemic disease associated with chorioretinitis

COMPLICATIONS

• Retinal hemorrhage/detachment, visual impairment with macular involvement

SIGNS & SYMPTOMS

 Floaters (vitritis), blurred vision, impaired color/night vision, ocular pain, photophobia, excessive lacrimation

DIAGNOSIS

DIAGNOSTIC IMAGING

Fluorescein angiography

Irregularities

Fundoscopy

 Creamy white/yellow/gray lesions; keratic precipitates; retinal edema, necrosis; chorioretinal atrophy, neovascularization; cotton-wool infiltrates (Candida-associated chorioretinitis); polymorphic retinochoroidal scars (toxoplasmosis-associated chorioretinitis)

OTHER DIAGNOSTICS

Clinical history, physical examination

TREATMENT

MEDICATIONS

Corticosteroids/antimicrobials



Figure 76.3 A retinal photograph displaying the features of chorioretinitis. There are numerous, patchy, cream-colored lesions and retinal edema.

CONJUNCTIVITIS

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PATHOLOGY & CAUSES

- Inflammation of conjunctiva, transparent mucous membrane covering inside of eyelids (tarsal conjunctiva), globe (bulbar conjunctiva)
 - Non-keratinized epithelium containing goblet cells, highly vascularized substantia propria
 - Turns pink/red when inflamed: diffuse conjunctival injection
- Infection, inflammation → dilatation of conjunctival vessels \rightarrow conjunctival hyperemia, edema → inflammatory discharge

TYPES

Infectious (bacterial)

Highly contagious; spread by direct contact

- Common causes: Staphylococcus aureus, Streptococcus pneumoniae, Haemophilus influenzae
- Hyperacute bacterial conjunctivitis
 - Causes: Neisseria gonorrhoeae (most common)/Neisseria meningitidis
 - Oculogenital disease: usually transmitted from genitals to eyes via hands
 - Vision-threatening
- Chlamydial
 - Caused by Chlamydia trachomatis
 - Adult inclusion conjunctivitis: chronic, indolent
 - Trachoma: infectious blindness cause worldwide; active trachoma caused by serotypes A, B, Ba, C (low-income country-endemic, mostly in children); initial follicular inflammation progresses in severity → cicatricial disease, vision

Infectious (viral)

- Highly contagious; spread by direct contact
- Causes: adenovirus (most common), HSV (in children), varicella zoster virus (VZV)
 - Ocular manifestation of systemic infection
 - Epidemic keratoconjunctivitis (EKC): caused by adenovirus 8, 19, 37; fulminant conjunctivitis, keratitis (epithelium of conjunctiva, cornea); corneal inclusions degrade visual acuity

Noninfectious (allergic)

- Caused by airborne allergens (seasonal, perennial)
- Immunoglobulin E (IgE)-mediated → local mast cell degranulation

Noninfectious (nonallergic)

Caused by mechanical/chemical insult

RISK FACTORS

- Exposure to causative agent, immunocompromised state, atopy (allergic conjunctivitis)
- Contact lens wear: common source of mechanical injury, nonallergic, infectious coniunctivitis

COMPLICATIONS

- Cornea: keratitis (inflammation), ulcer, perforation, scarring
- Dacryocystitis (bacterial infection of lacrimal sac)
- Vision loss

SIGNS & SYMPTOMS

- Appearance: unilateral/bilateral inflammation; pinkish-red eye; eyelid edema; chemosis (conjunctival edema); excessive lacrimation
- Discharge
 - Bacterial: purulent/mucopurulent; white/ yellow/green
 - Gonococcal: hyper-purulent, profuse
 - Viral: watery; stringy
 - Allergic: watery, mucoid
 - Nonallergic: mucoid

- Infected eye "stuck" shut from morning crusting; gritty, burning sensation (viral); itching (allergic); photophobia (corneal involvement); transient visual impairment
- Preauricular lymphadenopathy



Figure 76.4 The clinical appearance of conjunctivitis.

DIAGNOSIS

LAB RESULTS

- Adenoviral conjunctivitis: rapid point-ofcare adenovirus antigen test
- Recalcitrant conjunctivitis: conjunctival biopsy (rule out neoplasm)

Giemsa/gram stains

 Confirm identity of organism in suspected infectious cause

OTHER DIAGNOSTICS

Clinical history, physical examination

TREATMENT

MEDICATIONS

- Ocular lubricant drops/ophthalmic ointment
- Allergic conjunctivitis: antihistamine drops
- Adult inclusion conjunctivitis: systemic therapy to eradicate Chlamydia infection (antibiotics)
- Bacterial conjunctivitis: Topical antibiotic drops/ointment
- Epidemic keratoconjunctivitis (EKC): topical glucocorticoids

OTHER INTERVENTIONS

- Warm, wet compresses encourages drainage
- Hyperacute conjunctivitis, EKC: immediate specialized ophthalmologist referral
- Viral conjunctivitis: self-limiting; usually resolves in 2-3 weeks

KERATITIS

osms.it/keratitis

PATHOLOGY & CAUSES

- Cornea inflammation → corneal tissue destruction
- Inflammatory response → stromal damage from infection, host response \rightarrow edema, infiltrates, necrotic ulceration, focal thinning, perforation

CAUSES

Infectious

- Bacteria: Staphylococcus aureus, Pseudomonas aeruginosa, coagulasenegative Staphylococcus, diphtheroids, Streptococcus pneumoniae
- Viruses: HSV, herpes zoster
- Fungi: Candida supp., Aspergillus supp., Fusarium supp.
- Parasites: Acanthamoeba

Noninfectious

 Corneal inflammation with no known infectious etiology

RISK FACTORS

- Corneal epithelium disruption
 - Contact lenses (contact lens-related keratitis); esp. improper use (e.g. overnight wear, poor hygiene)
 - Recent keratoplasty, trauma, corneal exposure (e.g. Graves' ophthalmopathy, Bell's palsy)

- Immunocompromised state
- Topical (ocular) corticosteroid use
- Contributing disorders: rosacea; keratoconjunctivitis sicca (dry eye syndrome); neurotrophic keratitis (lesion on cranial nerve V); autoimmune diseases (e.g. rheumatoid arthritis, cicatricial pemphigoid)

COMPLICATIONS

• Endophthalmitis (interior eye inflammation), intraocular damage, vision loss, keratolysis (corneal melting)

SIGNS & SYMPTOMS

- Erythema
- Preauricular lymphadenopathy
- Discharge: mucopurulent (bacterial), watery (viral)
- Corneal opacity, stromal infiltrate (immune complex deposits), ulcer
 - Bacterial keratitis: yellow infiltrates
 - Fungal keratitis: white infiltrates, feathery borders
 - Acanthamoeba: Wessely ring infiltrate
- Hypopyon (layer of white cells in anterior chamber): fulminant bacteria
- Foreign body sensation; difficulty keeping eye open; photophobia; pain; decreased visual acuity, blurred vision; blepharospasm



Figure 76.5 An individual with sterile keratitis of the left eye.

DIAGNOSIS

DIAGNOSTIC IMAGING

Fundoscopy

 Slit beam; examine contour abnormalities of cornea, lens, retina; small corneal infiltrates; faint branching grey opacity (viral keratitis)

LAB RESULTS

 Corneal scrapings, cultures: suspected infectious etiology

OTHER DIAGNOSTICS

Clinical history, physical examination

Penlight

 Visualizes infiltrate/ulcer (> 0.5mm); round, white spot (bacterial keratitis)

Fluorescein dye

- Corneal uptake of dye
 - Visualize loss of epithelial cells, ulceration
 - Green glow under cobalt blue light
 - Diffuse white opacity/dull corneal light reflex
 - Seidel sign (leaking aqueous humor → fluorescein streaming): penetrating trauma

Snellen chart

J visual acuity

TREATMENT

MEDICATIONS

Topical antimicrobials for infectious etiology

OTHER INTERVENTIONS

- Control of associated comorbidities
- Temporary discontinuation of wearing contact lenses

ORBITAL CELLULITIS

osms.it/orbital-cellulitis

PATHOLOGY & CAUSES

 Serious infection involving contents of orbit (ocular muscles, surrounding fat; not globe)

CAUSES

- Entry of microorganisms into orbital space
 - Via anatomical perforations of nerves, blood vessels in paranasal sinuses (e.g.
 - Migration from surrounding tissues (e.g. face, eyelids) after local trauma/surgery
 - □ Inflammatory response → tissue destruction

RISK FACTORS

- More common in children
- Migration from other infections
 - Bacterial rhinosinusitis: Staphylococcus aureus, streptococci (common); fungal rhinosinusitis (rare)
 - Dacryocystitis: lacrimal sac infection
 - Infected mucocele: mucus-containing cystic lesion of salivary gland
 - Infections involving teeth, middle ear,
- Direct inoculation: ophthalmic surgical procedures; orbital trauma with fracture/ foreign body

COMPLICATIONS

- Extraorbital extension: epidural/subdural empyema; brain abscess; meningitis; cavernous sinus thrombosis; dural sinus thrombosis: involvement of cranial nerves III, IV, V, VI; optic neuritis
- Endophthalmitis: interior eye inflammation
- Vision loss
- Potentially fatal if sepsis develops

SIGNS & SYMPTOMS

Systemic

 Fever; severe headache, vomiting, mental status changes (intracranial complications)

Ocular

• Red, swollen eyelids; chemosis (conjunctival edema); pain (esp. with eye movement); ophthalmoplegia (paralysis of eye muscles); proptosis (abnormal displacement of eye); impaired visual acuity, color vision; abnormal pupillary light reflex

DIAGNOSIS

DIAGNOSTIC IMAGING

CT scan/MRI

• Orbits, sinuses; detects abscess, intracranial changes

Dilated fundoscopy

 Determines optic neuropathy/retinal vascular occlusion

LAB RESULTS

Complete blood count (CBC)

 Leukocytosis; ↑ absolute neutrophil count (ANC)

Blood/orbital/subperiosteal aspirates cul-

Identify causative organism

OTHER DIAGNOSTICS

- Clinical history, physical examination
- Ocular motility: pain with movement
- Pupillary light reflex: sluggish/absent reflex → optic nerve involvement
- Exophthalmometry: measures degree of proptosis
- Asses color vision acuity: determines optic nerve involvement
- Intraocular pressure measurement (1)

TREATMENT

MEDICATIONS

Antimicrobials

SURGERY

 External (through orbit)/endoscopic transcaruncular approach

CHARACTERISTICS OF PERIORBITAL & ORBITAL CELLULITIS

	PERIORBITAL CELLULITIS	ORBITAL CELLULITIS
EYE SWELLING WITH/WITHOUT ERYTHEMA	+	+
FOCAL PAIN	+/-	+
PAIN WITH MOVEMENT	-	+
PROPTOSIS	-	+
OPHTHALMOPLEGIA	-	+
DECREASED VISUAL ACUITY	-	+
CHEMOSIS	-	+/-
IMPAIRED PUPILLARY FUNCTION	-	+
FEVER	+/-	+
LEUKOCYTOSIS	+/-	+/-
POTENTIAL FOR VISION LOSS	-	+

PERIORBITAL (PRESEPTAL) CELLULITIS

osms.it/periorbital-cellulitis

PATHOLOGY & CAUSES

• Mild infection of superficial tissues of anterior eyelid (tissues anterior to orbital septum): more common than orbital cellulitis

CAUSES

Introduction/migration of microorganisms into preseptal space: Staphylococcus aureus, Streptococcus pneumoniae, other streptococci, anaerobes

RISK FACTORS

- More common in children
- Migration from other infections: sinusitis: upper respiratory tract infection; dacryocystitis; bacteremia (rare)
- Direct inoculation: trauma (e.g. insect bites, animal bites, introduction of foreign bodies); ophthalmic surgical procedures

COMPLICATIONS

Orbital cellulitis



Figure 76.6 An individual with left-sided periorbital cellulitis.

SIGNS & SYMPTOMS

• Ocular pain, eyelid swelling, erythema, fever, lymphadenopathy

DIAGNOSIS

DIAGNOSTIC IMAGING

Contrast-enhanced CT scan (orbits, sinuses)

 Distinguishes between preseptal, orbital cellulitis; associated sinusitis

LAB RESULTS

CBC

Leukocytosis

Cultures (abscess contents, paranasal sinus secretions)

• Identify causative agent

OTHER DIAGNOSTICS

Clinical history, physical examination

TREATMENT

MEDICATIONS

Oral antibiotics

STYE (HORDEOLUM)

osms.it/stye

PATHOLOGY & CAUSES

 Blockage, purulent inflammation of upper/ lower evelid

CAUSES

 Sterile/bacterial (e.g. Staphylococcus aureus, Staphylococcus epidermidis)

• Meibomian sebaceous gland; points toward conjunctival side of lid \rightarrow conjunctival inflammation

External

 Zeiss/Moll sebaceous glands; points toward skin surface of eyelid

RISK FACTORS

 Touching eyes with contaminated hands, chronic blepharitis, seborrhea, improper contact lens hygiene, sleeping with eye makeup, immunocompromised state

COMPLICATIONS

Hardens → chalazion



Figure 76.7 A stye on the right lower eye

SIGNS & SYMPTOMS

 Tenderness; fluctuant pustule; localized swelling, erythema; excessive lacrimation; photophobia

DIAGNOSIS

DIAGNOSTIC IMAGING

Slit lamp, fundoscopy

• Determine infection extension to other tissues

OTHER DIAGNOSTICS

- Clinical history, physical examination
- Visual acuity assessment

TREATMENT

MEDICATIONS

• Topical antibiotic ointment

SURGERY

• Incision, curettage: if progresses to chalazion

OTHER INTERVENTIONS

- Warm compresses encourage drainage
- Usually self-limiting with spontaneous resolution

EYELID LESIONS OVERVIEW		
	STYE (HORDEOLUM)	CHALAZION
ONSET	Acute	Chronic
CAUSE	Usually infectious	Noninfectious
GLAND INVOLVEMENT	Internal stye - Involves meibomian gland - Lesion points toward conjunctival side of lid External stye - Involves Zeiss/Moll glands - Lesion points toward skin surface of eyelid	Deep chalazion - Involves meibomian gland Superficial chalazion - Involves Zeis glands Lesions are deeper than hordeolum
PAIN	Yes	No; possible mild tenderness
INFLAMMATORY CHARACTERISTICS	Purulent	Lipogranulomatous
LESION CHARACTERISTICS	Small, fluctuant	Large, firm, rubbery, nodular
DURATION OF SYMPTOMS	Self-limiting with spontaneous drainage	May wax, wane; often lasts for months
SEVERE COMPLICATION	May evolve into chalazion if healing compromised	May be sign of eyelid carcinoma

UVEITIS

osms.it/uveitis

PATHOLOGY & CAUSES

- Inflammation of uveal tract (choroid, ciliary body, iris); unilateral/bilateral
- Onset: rapid/insidious
- Course: acute/recurrent/chronic
- Duration: persistent (> three months)/ limited (≤ three months)

TYPES

Anterior (most common)

 Anterior uveal tract; iritis, iridocyclitis (inflammation of ciliary body)

Panuveitis

 Anterior chamber, vitreous body, retina/ choroid

Posterior uveitis

Retina/choroid

Intermediate uveitis

Vitreous body; chorioretinal inflammation

CAUSES

- Bacterial: tuberculosis, syphilis
- Viral: cytomegalovirus, HSV
- Fungal: candidiasis, Pneumocystis jirovecii
- Parasitic: Acanthamoeba, toxoplasmosis
- Noninfectious systemic: Crohn's disease, ankylosing spondylitis
- Conditions confined to eye: trauma, acute retinal necrosis

RISK FACTORS

 Systemic infectious, inflammatory conditions

COMPLICATIONS

 Intraocular hypertension, glaucoma; increased intraocular pressure; posterior synechiae (iris adheres to lens); band keratopathy (corneal calcium deposits); cataract: vision loss

SIGNS & SYMPTOMS

- Ocular erythema
- Impaired vision
- Pain, photophobia, vision distortion, floaters (vitritis), photopsia (flashing lights)

DIAGNOSIS

DIAGNOSTIC IMAGING

Fluorescein/indocyanine green angiography (posterior uveitis)

 Evaluate status of retinal vascular circulation; identify choroidal disease

Fundoscopy

- Ciliary flush: perilimbal redness
- Keratic precipitates: inflammatory deposits on cornea
- Hypopyon: white blood cells settle on bottom of anterior chamber
- Haziness of aqueous humor: protein accumulation

LAB RESULTS

Microscopy, cytology, culture, polymerase chain reaction (PCR)

• Fluid sampling/biopsy; identify presence of infectious agent

OTHER DIAGNOSTICS

Clinical history, physical examination

Snellen chart

J visual acuity

Pupillary light reflex

 Sluggish pupillary reaction to light → synechiae

Intraocular pressure

 No change if uncomplicated uveitis; ↑ in acute uveitis-induced glaucoma

TREATMENT

MEDICATIONS

- Corticosteroids: topical, local injection, implantable, systemic
- Recalcitrant uveitis: immunomodulatory agents (if corticosteroid response inadequate)
- Recalcitrant uveitis: tumor necrosis factor (TNF) inhibitor (if resistant to treatment)
- Posterior synechiae prevention: mydriatic/ cycloplegic medications
- Viral-associated uveitis: antivirals



Figure 76.8 An individual with a hypopyon of the left eye as a result of severe anterior uveitis.