Introduction

The red eye is the most common ocular complaint in patients seeking medical assistance at primary healthcare level. The differential diagnosis of the red eye varies from relatively innocuous and trivial conditions to those that are more devastating and potentially blinding. The purpose of this article is to help identify certain key clinical features of the discussed conditions, to help avoid pitfalls in diagnosis and to convey appropriate treatment modalities.

Conjunctivitis

Conjunctivitis is defined as inflammation of the conjunctival membrane that covers the ocular surface. Potential causes are bacterial, viral and allergic.

Viral conjunctivitis

The most common responsible viruses include adenovirus, herpes simplex virus and molluscum contagiosum. Typically, viral infection is characterised by an acute follicular conjunctival reaction (Figure 1), as well as preauricular or submandibular lymphadenopathy.

Adenovirus conjunctivitis

The severity of adenoviral conjunctivitis may vary from a mild infection to more severe syndromes, such as pharyngoconjunctival fever and epidemic keratoconjunctivitis. Both of these are highly contagious and tend to occur in epidemics. Pharyngoconjunctival fever affects children predominantly and is often accompanied by an upper respiratory tract infection. Epidemic keratoconjunctivitis ("pink eye") is not typically associated with systemic manifestations. These infections induce an acute follicular conjunctival reaction which is often bilateral and associated with preauricular lymphadenopathy. The incubation period is usually 7-9 days before the onset of symptoms. Patients report ocular itching, tearing, redness and photophobia. In severe cases, subconjunctival bleeding may occur and the cornea may become affected in the second week.

Treatment of adenoviral conjunctivitis is supportive mainly. Patients should be instructed to use cold compresses and lubricating eye drops for comfort. Topical antihistamine eye drops, such as Spersallerg®, may be used to alleviate severe itching.

Herpes simplex conjunctivitis

Primary ocular herpes simplex virus infection affects infants and young children predominantly. Patients may present with a follicular conjunctivitis that manifests as a red, teary eye, associated with typical vesicular eruption on the eyelids (Figure 2). Treatment is the same as that for herpes simplex keratitis. The treatment of skin lesions may also include the topical administration of acyclovir ointment.
Molluscum contagiosum keratoconjunctivitis

Molluscum contagiosum can cause follicular conjunctivitis in association with an eyelid lesion. Usually, the lesion is a small, pearly umbilicated nodule on or near the lid margin (Figure 3). Multiple lesions may be present, especially in a patient with human immunodeficiency virus (HIV). Treatment involves incision and curettage of the symptomatic lesions. Asymptomatic lesions, if left long enough, are usually self-limiting.

Bacterial conjunctivitis

Acute bacterial conjunctivitis

Acute bacterial conjunctivitis is a very common ocular condition which is primarily due to the *Staphylococcus*, *Haemophilus* and *Streptococcus* species. These organisms may be spread by hand-to-eye contact or by colonisation of adjacent mucosal tissues, such as the nasal or sinus mucosa.

Initially, acute bacterial conjunctivitis presents unilaterally. The second eye is often affected soon thereafter. Symptoms include a sticky discharge that causes the eyelids to adhere to each other in the morning and a foreign body sensation. Usually, examination reveals generalised injection of the conjunctiva (Figure 4). Vision and pupillary reactions should be normal. The cornea and anterior chamber should be clear. Generally, bacterial conjunctivitis is benign and self-limiting in nature and is highly responsive to empirical therapy. It is also a contagious condition so patients should be instructed in proper hygiene and hand washing. Chloramphenicol, in the form of ointment or drops, is a good initial empirical treatment as it has broad-spectrum cover. Patients should be referred if the diagnosis is unclear or if the conjunctivitis is recurrent or refractory to initial therapy.

Gonococcal conjunctivitis

Typically, this ocular disease presents as a hyperacute purulent conjunctivitis. A profuse purulent discharge is often present and is associated with periorbital oedema, marked conjunctival hyperaemia and chemosis (severe conjunctival swelling) (Figure 5), as well as preauricular lymphadenopathy.

Prompt treatment of gonococcal conjunctivitis is essential since this organism can rapidly cause corneal ulceration and perforation. Ideally, because of the rapid progression of this condition, patients should be referred to an eye specialist after commencing initial management with saline irrigation to clear the purulent material from the lids and conjunctiva. Hourly topical fluoroquinolone drops (ofloxacin or ciprofloxacin) should be given, as well as an
intramuscular loading dose of ceftriaxone 1 g. Oral treatment with either erythromycin 250-500 mg four times a day, or doxycycline 100 mg twice daily for possible coexisting chlamydial infection, is also advised.

**Chlamydial conjunctivitis**

Chlamydia is a common cause of chronic conjunctivitis and is responsible for the two clinical entities of adult inclusion conjunctivitis and trachoma.

- **Adult inclusion conjunctivitis**

  This is a sexually transmitted infection that typically affects young, sexually active adults. It presents as a bilateral follicular conjunctivitis with a mucopurulent discharge and assumes a chronic course if left untreated. Non-specific urethritis or cervicitis is commonly associated with this condition. Management involves oral administration of doxycycline 100 mg twice daily for three weeks, as well as screening for HIV and syphilis. Erythromycin is a good alternative in patients who are allergic to tetracyclines. All sexual partners should also be evaluated and treated accordingly.

- **Trachoma**

  Trachoma is one of the most common causes of preventable blindness in the world. It is extremely common in equatorial Africa. It occurs predominantly in low socio-economic settings in association with poor sanitation and hygienic conditions. The house fly is responsible for the infection-reinfection cycle.

  The initial conjunctival inflammation is called “active trachoma”, and is predominantly seen in children, especially preschool children. It is characterised by mucopurulent conjunctivitis. The conjunctival surface of the upper eyelid shows follicles that may also appear at the junction of the cornea and the sclera (limbal follicles). With healing, these limbal follicles leave small depressions called Herbert’s pits. These are characteristic of trachoma.

  The latter structural changes of trachoma are referred to as “cicatricial trachoma”. These include scarring of the palpebral conjunctiva, as well as the formation of fibrovascular pannus (blood vessels and scar tissue that invade the superior cornea). The cicatricial phase of the disease results in entropion (in-turned eyelids) and trichiasis (in-turned eyelashes) (Figure 6), and is responsible for the blinding complications of corneal ulceration, scarring and opacification.

  National governments implement trachoma control programmes using the World Health Organization’s recommended SAFE strategy, which includes:

  - **S**urgery to correct the advanced stages of the disease.
  - **A**ntibiotics, such as doxycycline or azithromycin, to treat the active stage of the disease.
  - **F**acial washing to reduce disease transmission.
  - **E**nvironmental changes to improve access to clean water and sanitation.

Figure 6: Trichiasis (in-turned eyelashes) are associated with trachoma

**Allergic conjunctivitis**

**Acute allergic rhinoconjunctivitis**

Acute allergic rhinoconjunctivitis (hayfever) is the most common form of ocular and nasal allergy.

Presentation is with transient attacks of itchiness, teariness, and redness associated with sneezing, as well as nasal discharge. Lid oedema and chemosis may also be present in severe cases. Treatment involves the use of antihistamine drops to provide symptomatic relief and mast cell stabilisers, e.g. sodium chromoglycate, four times a day for long-term use.

**Vernal keratoconjunctivitis**

Vernal keratoconjunctivitis is a bilateral, recurrent disorder that is very common in the drier parts of South Africa. It affects children and teenagers predominantly and tends to worsen seasonally. There is often associated atopy with a history of asthma and eczema in infancy.

Characteristic symptoms include intense itching associated with tearing, photophobia and the sensation of a foreign body in the eye. Secondary skin changes of the eyelids are a common consequence of persistent eye rubbing. Traditionally, this condition is classified as either a palpebral, limbal or mixed type.

Palpebral disease is typified by the presence of diffuse papillary hypertrophy of the superior tarsal conjunctiva, associated with thick mucoid deposition between the papillae (Figure 7). Limbal disease is characterised by the presence of gelatinous papillae on the limbal conjunctiva (Figure 8). In severe cases, the cornea may also be affected with erosions and dry white plaques called shield ulcers (Figure 9). Treatment may be complex. Most patients require ophthalmic assessment. Mild cases require the use of a mast cell stabiliser, which is the cornerstone of treatment, and antihistamine drops. Topical steroids are effective, but should only be given under medical supervision.

**Keratitis**

Keratitis may be noninfective or infective in origin. Infective keratitis is often a unilateral condition and may be caused by a variety of organisms, including bacterial, viral and fungal...
elements. Usually, the symptoms of a red eye associated with pain, photophobia, decreased vision and a discharge occur.

**Bacterial keratitis**

Bacterial keratitis is uncommon in a normal eye as the corneal epithelium provides a barrier against many organisms. Most central corneal ulcers follow a breach in this layer, secondary to either trauma or pre-existing ocular surface disease. Other risk factors include a history of wearing contact lens and systemic immunosuppression. The most common implicated organisms are *Pseudomonas*, *Staphylococcus* and *Streptococcus*.

Signs include an epithelial defect which stains with fluorescein, stromal infiltrates and a hypopyon (pus in the anterior chamber) (Figure 10). Initial treatment in a primary care setting may include discontinuation of wearing contact lenses and the use of a plastic eye shield for protection. Since bacterial keratitis has the potential to progress rapidly to corneal perforation, these patients need prompt referral to specialist intervention. Intensive topical antibiotics and atropine drops are given to sterilise the ulcer and relieve associated ciliary spasms respectively.

**Herpes simplex keratitis**

Herpetic eye disease represents one of the major causes of unilateral corneal scarring worldwide. As much as 60% of corneal ulcers may be the result of herpes simplex virus. Typically, signs include decreased visual acuity, a watery discharge and single, or less commonly, multiple, linear branching epithelial lesions called dendritic ulcers. Usually, corneal sensation is reduced and the ulcer bed stains with fluorescein (Figure 11).

Acyclovir ointment should be applied five times daily for the first few days. Recurrences are not uncommon so the treatment may need to be repeated. At all costs, topical steroids should be avoided in these patients as corneal perforation may ensue.

**Herpes zoster ophthalmicus**

Herpes zoster ophthalmicus is a painful vesicular skin rash, caused by reactivation of the varicella-zoster virus which occurs unilaterally over the forehead. Diagnosis in a young patient suggests the possibility of underlying HIV infection or immunosuppression. Typical skin eruption is obvious, but a valuable indicator of potential ocular involvement is the presence of skin lesions that affect the tip of the nose. This is referred to as Hutchinson’s sign (Figure 12).

Management consists of local wound care, systemic acyclovir in high doses and early referral. Oral acyclovir has been shown to shorten the duration of signs and symptoms and appears to be most beneficial if instituted within 72 hours of the onset of skin lesions.
Fungal keratitis

Fungal keratitis remains a significant cause of blindness in the developing world. Local predisposing factors include trauma, especially with vegetable or organic matter, contact lenses and topical steroids.

Usually, fungal ulcers are much less aggressive than bacterial ulcers. Typically, presentation is a red, painful eye associated with a decrease in visual acuity. These ulcers are slow-spreading and are dull grey in appearance, with feather-like extensions. Generally, satellite lesions and a hypopyon are also evident (Figure 13).

Successful resolution of the disease depends on a high degree of suspicion and early institution of therapy. Thus, immediate referral of these patients is necessary. Management consists of topical and systemic antifungals and cycloplegic drops, as well as treating coexistent bacterial infection that is often present.

Acute angle closure

Acute angle closure is a condition in which the intraocular pressure rises rapidly to very high levels. It represents an ophthalmic emergency and requires immediate referral to specialist intervention.

Acute angle closure classically occurs in older, far-sighted patients. Symptoms include severe unilateral eye pain associated with a headache on the same side as the painful eye, decreased vision and marked redness of the eye. Constitutional symptoms often include nausea and vomiting. An examination reveals diffuse conjunctival injection, a hazy cornea, a shallow anterior chamber, high intraocular pressure and a fixed, mid-dilated pupil (Figure 14). The aim of initial treatment is to lower the pressure as quickly as possible with a loading dose of 500 mg acetazolomide taken orally. This can be initiated in a primary care setting before transferring the patient if one is highly suspicious of the diagnosis, provided the patient is not allergic to sulphonamides. Antiemetics and analgesia should be given as necessary.

Anterior uveitis

Uveitis is defined as inflammation that affects one or more parts of the uveal tract. The uvea is the vascular coat of the eye and consists of the iris, ciliary body and the choroid. Anterior uveitis is the most common form of uveitis and predominantly affects the iris (iritis).2

Usually, presentation occurs with the acute onset of unilateral pain, redness, lacrimation, photophobia and a moderate reduction in visual acuity. The redness is often more marked in the circumcorneal region. The pupil is miotic and irregular due to
adhesions that form between the pupil margin and the anterior surface of the lens (posterior synechiae) (Figure 15).

Clusters of cells on the corneal endothelium, called keratic precipitates (Figure 16), may also be seen, and if the uveitis is severe, a hypopyon may form. Diagnosis is dependent on slit-lamp examination of the eye so most patients require referral. Initially, topical steroids are used frequently, then tapered over several days to weeks, depending on the severity of the inflammation. Adequate cycloplegia using atropine 1% is maintained during the course of the treatment to relieve pain and also to dilate the pupil which prevents the formation of posterior synechiae.

Penetrating trauma

Penetrating injuries to the eye are serious as they are potentially blinding. However, life-threatening conditions should be recognised and immediately treated as these take precedence over any associated ocular injuries. Most of these injuries are obvious, although a high index of suspicion is needed to diagnose an occult injury. Signs suggestive of this type of penetrating injury include severe conjunctival swelling, an anterior chamber that looks deeper than the other eye, and a soft eye.

These patients require urgent referral as surgical repair is mandatory. Manipulation of an open globe can result in exacerbation of the initial injury, as even the smallest amount of pressure placed on the eye can lead to extrusion of the ocular contents, and thus should be avoided. The eye should be covered with a protective plastic shield, not an eye pad, and use of topical medication avoided. If a plastic shield is not available, the bottom half of a clean polystyrene cup can be used instead. A dose of oral ciprofloxacin should be given before the patient is referred. This is to attempt to prevent infection of the ocular contents (endophthalmitis).

Burn wounds

Chemical burns

Chemical burns, especially those caused by alkali, constitute an ophthalmic emergency. Prompt irrigation of the eye is most important to limit the extent of damage from such burns. A few litres of sterile isotonic saline should be rinsed through the palpebral opening with the aid of an eyelid speculum and topical anaesthetic to prevent reflex blepharospasm. Complete removal of the offending agents, by sweeping under the eyelids with a wet cotton bud, should also be attempted. The end-point of this initial management is to achieve a pH of between 7.3-7.7 measured with the appropriate block on a urine dipstick. Systemic analgesia, as well as topical dilating drops, also induce comfort. The patient should be referred for further evaluation by an ophthalmologist.

Thermal burns

Usually, these injuries are less severe than their chemical counterparts. Topical antibiotics, dilating drops, as well as systemic analgesics, constitute a good initial empirical approach. The patient should be referred for further evaluation and treatment of possible long-term complications.

Conclusion

This article has outlined a wide variety of conditions with which patients may present. A red eye is implicated in all of them. If the diagnosis is approached systematically by taking a thorough history from the patient and knowing what to look for during the examination, in many cases, it is possible to make the correct diagnosis. Once a diagnosis has been made, it is very important to know which conditions may be safely treated at primary care level and which require urgent referral for more specialised management.

Bibliography