Ofloxacin 0.3% eye drops

Alert	Ofloxacin eye drops are not recommended for empirical treatment of bacterial conjunctivitis in neonates.
	Use under close supervision and in consultation with an ophthalmologist.
Indication	Treatment of external bacterial eye infections including keratitis and bacterial conjunctivitis.
Action	Bactericidal by inhibiting bacterial DNA synthesis by blocking DNA gyrase and topoisomerase IV.
Drug type	Broad spectrum fluoroquinolone antibiotic
Trade name	Ocuflox
Presentation	0.3% (3 mg/mL ofloxacin), 5 mL dropper bottle
Dose	Dose frequency depends on severity of infection and response to treatment ^{1,2}
	First 48 hours: 1 drop every 2-4 hours in the affected eye and, if clinical improvement,
	From Day 3 and up to day 10: 1 drop every 6 hours.
	May require more frequently (e.g. every 1 hour) in severe infections.
Dose adjustment	Therapeutic hypothermia – Not applicable
	ECMO – Not applicable
	Renal impairment – Not applicable
	Hepatic impairment – Not applicable
Maximum dose	
Total cumulative	
dose	
Route	Topical
Preparation	Not required.
Administration	Instil 1 eye drop in the affected eye/s.
	After administering eye drop, gently press against the inner corner of eye to reduce systemic absorption.
	If other eye drop(s) are administered, wait for 5 minutes between drops.
Monitoring	
Contraindications	History of hypersensitivity with quinolone useor any other component of the formulation.
Precautions	
Drug interactions	
Adverse reactions	Mild transient ocular irritation, white corneal precipitates, keratitis, allergic reactions {hypersensitivity
	(very rare) including angioedema, anaphylaxis and Stevens-Johnson syndrome}
Compatibility	Not applicable
Incompatibility	Not applicable
Stability	
Storage	Store below 25°C. Protect from light and excessive heat.
	To avoid contamination of the solution, keep container tightly closed.
	Discard container 4 weeks after opening.
Excipients	Ocuflox eye drops contain 0.05 mg/mL benzalkonium chloride.
Special comments	

Newborn use only

Topical 0 3% ofloxacin is an effective and safe treatment for conjunctivitis and blepharoconjunctivitis in paediatric and adult patients. Data in neonates are very limited.
paediatric and adult patients. Data in neonates are very limited.
Bron et al compared a 7 day course of 0.3% ofloxacin with chloramphenicol 0.5% in 167 adult patients
with suspected bacterial external ocular infection. Patients were instructed to use their eye drops every
2-4 waking hours (according to severity as determined by the physician) on days 1 and 2 and 4 times date
from days 3 to 7. In the study, 63% patients had conjunctivitis, while blepharoconjunctivitis,
Keratoconjunctivitis and blepharitis were present in 20%, 7% and 7% respectively. Microbiological
improvement rates were 85% and 88% in ofloxacin and chloramphenicol groups respectively. Bacteria
were eradicated in 79% and 77% of ofloxacin and chloramphenicol groups respectively. Clinical
improvement rates were 100% for ofloxacin, and 95% for chloramphenicol.
In a randomised controlled trial, Gwon et al administered topical 0.3% of loxacin in 93 patients with
suspected bacterial external ocular infection. One drop of onoxacin was applied to the affected eye(s) si
(day 2 to day 10) The control group received 0.3% contamicin in a similar decage schedule. Paseline
(udy 5 to udy 10). The control group received 0.5% gental inclining similar dosage schedule. Baseline
gentamicin group. Among patients treated with ofloyacin 98% (51/52) were either clinically cured or
improved by day 11, compared with 92% (48/52) of the gentamicin group. Microbiological improvemen
was achieved in 78% (40/51) of the ofloxacin patients, compared with 67% (35/52) of the gentamicin
group. Bacterial proliferation occurred in 16% (8/51) of the ofloxacin group vs 27% (14/52) of gentamici
treated subjects. ² Suggested dose regimens in this formulary are extrapolated from the above 2 studies.
^(1,2) . However, one study found that administration of topical Ofloxacin 0.3% solution 12 hourly was as
efficacious as 6 hourly schedule in clinical and bacteriological improvement in patients with conjunctivit
and blepharoconjunctivitis. ⁵
Safety
Bron et al noted no difference in adverse reactions between ofloxacin and chloramphenicol groups. Of 8
patients receiving ofloxacin, one participant developed haemorrhagic conjunctivitis and marked
palpebral oedema. ¹ In the RCT by Gwon et al, adverse reactions possibly caused by drug treatment were
included huming, stinging, and photophobic possisiating discontinuation of the drug. No drug
treatment related effects on visual acuity, onbthalmoscony findings, or lens nathology were observed
There was no notable difference between treatment groups in comfort of drug application. In one study
ofloxacin 0.3% or azithromycin eve drops were administered for 7 days for postoperative
endophthalmitis prophylaxis after intravitreal injections. Five patients in the ofloxacin group and two
patients in the Azithromycin group developed endophthalmitis following intravitreal Ranibizumab
injection (p<0.05). ³ In a case report, Claerhout reported corneal deposits in two children with
keratoconjunctivitis who received 0.3% ofloxacin. ⁴
Practice points Due to concern about emerging resistance:
Reserve quinolones for treatment of bacterial keratitis (under close supervision by, or following
discussion with, an ophthalmologist)
Other antibacterials are preferred for empirical treatment of conjunctivitis.
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