1. NAME OF THE MEDICINAL PRODUCT

Tobramycin 0.3%

2. QUALITATIVE AND QUANTITATIVE COMPOSITION

Each ml contains: tobramycin 3 mg

3. PHARMACEUTICAL FORM

Eye drops, suspension

4. CLINICAL PARTICULARS

4.1 Therapeutic indications

Prevention and treatment of inflammation and prevention of infection associated with cataract surgery in adults and children aged 2 years and older.

4.2 Posology and method of administration

Posology

Adults

One drop instilled into the conjunctival sac(s) every 4 to 6 hours while the patient is awake. During the initial 24 to 48 hours, the dosage may be increased to one drop every two hours while the patient is awake. Dosing should continue for 14 days not to exceed a maximum of 24 days. Frequency should be decreased gradually as warranted by improvement in clinical signs. Care should be taken not to discontinue therapy prematurely.

Use in the Elderly

Clinical studies have indicated dosage modifications are not required for use in the elderly.

Paediatric population

Tobramycin may be used in children 2 years of age and older at the same dose as in adults. Currently available data is described in section 5.1.

The safety and efficacy in children younger than 2 years of age have not been established, and no data are available.

Use in hepatic and renal impairment

Tobramycin has not been studied in these patient populations.

Method of administration

Ocular use.

Shake the bottle well before use. To prevent contamination of the dropper tip and suspension, care should be taken not to touch the eyelids, surrounding areas, or other surfaces with the dropper tip of the bottle. Keep the bottle tightly closed when not in use. After cap is removed, if tamper evident snap collar is loose, remove before using product.

Gently closing the eyelid (s) and nasolacrimal occlusion for at least 1 minute after instillation is recommended. This may reduce the systemic absorption of medicinal products administered via the ocular route and result in a decrease in systemic side effects.

In case of concomitant therapy with other topical ophthalmic medicinal products, an interval of 5 minutes should be allowed between successive applications.

Eye ointments should be administered last.

4.3 Contraindications

- Hypersensitivity to Tobramycin
- Herpes simplex keratitis
- Vaccinia, varicella and other viral disease of the cornea and conjunctiva
- Mycobacterial infections of the eye caused by, but not limited to, acid-fast bacilli such as Mycobacterium tuberculosis, Mycobacterium leprae, or Mycobacterium avium.
- Fungal diseases of ocular structures or untreated parasitic eye infections.
- Untreated purulent infection of the eye.

4.4 Special warnings and precautions for use

Tobramycin is for topical use only and **not** for injection or oral use.

Prolonged use of topical ophthalmic corticosteroids (i.e. longer than the maximum duration used in clinical trials [24 days]) may result in ocular hypertension/glaucoma with resultant damage to the optic nerve and reduced visual acuity and visual fields defects and may also result in posterior subcapsular cataract formation.

Visual disturbance

Visual disturbance may be reported with systemic and topical corticosteroid use. If a patient presents with symptoms such as blurred vision or other visual disturbances, the patient should be considered for referral to an ophthalmologist for evaluation of possible causes which may include cataract, glaucoma or rare diseases such as central serous chorioretinopathy (CSCR) which have been reported after use of systemic and topical corticosteroids.

Prolonged use may also result in secondary ocular infections due to suppression of host response. Corticosteroids may reduce resistance to and aid in the establishment of bacterial, viral, fungal or parasitic infections and mask the clinical signs of infection.

Sensitivity to topically administered aminoglycosides may occur in some patients. Severity of hypersensitivity reactions may vary from local effects to generalized reactions such as erythema, itching, urticarial, skin rash, anaphylaxis, anaphylactoid reactions, or bullous reactions. If hypersensitivity develops during use of this medicine, treatment should be discontinued.

Cross-hypersensitivity to other aminoglycosides can occur, and the possibility that patients who become sensitized to topical tobramycin may also be sensitive to other topical and/or systemic aminoglycosides should be considered.

Serious adverse reactions including neurotoxicity, ototoxicity and nephrotoxicity have occurred in patients receiving systemic aminoglycoside therapy. Caution is advised when Tobramycin eye drops are used concomitantly with systemic aminoglycosides.

Caution should be exercised when prescribing Tobramycin eye drops to patients with known or suspected neuromuscular disorders such as myasthenia gravis or Parkinson's disease. Aminoglycosides may aggravate muscle weakness because of their potential effect on neuromuscular function.

Fungal infection should be suspected in patients with persistent corneal ulceration. If fungal infection occurs, corticosteroids therapy should be discontinued**.**

Prolonged use of antibiotics such as tobramycin may result in overgrowth of non-susceptible organisms, including fungi. If superinfection occurs, appropriate therapy should be initiated.

Topical ophthalmic corticosteroids may slow corneal wound healing. Topical NSAIDs are also known to slow or delay healing. Concomitant use of topical NSAIDs and topical steroids may increase the potential for healing problems. See section 4.5.

In those diseases causing thinning of the cornea or sclera, perforations have been known to occur with the use of topical corticosteroids.

4.5 Interaction with other medicinal products and other forms of interaction

No known drug interaction has been recorded of Artemether.

4.6 Pregnancy and lactation

Pregnancy

There are no or limited amount of data from the topical ocular use of tobramycin in pregnant women. Tobramycin does cross the placenta into the fetus after intravenous dosing in pregnant women. Tobramycin is not expected to cause ototoxicity from in utero exposure. Prolonged or repeated corticoid use during pregnancy has been associated with an increased risk of intra-uterine growth retardation. Infants born of mothers who have received substantial doses of corticosteroids during pregnancy should be observed carefully for signs of hypoadrenalism.

Breastfeeding

Tobramycin is excreted in human milk after systemic administration. It is unknown whether tobramycin is excreted in human milk following topical ocular administration. It is not likely that the amount of Tobramycin would be detectable in human milk or be capable of producing clinical effects in the infant following topical use of the product.

A risk to the suckling child cannot be excluded. A decision must be made whether to discontinue breast-feeding or to discontinue/abstain from therapy taking into account the benefit of breast feeding for the child and the benefit of therapy for the woman.

Fertility

Studies have not been performed to evaluate the effect of tobramycin on human or animal fertility.

4.7 Effects on ability to drive and use machines

None

4.8 Undesirable effects

Summary of the safety profile

In clinical studies involving over 1600 patients, Tobramycin was administered up to six times daily. No serious ophthalmic or systemic adverse reactions related to Tobramycin or components of the combination were reported in clinical studies. The most frequently reported adverse reactions with Tobramycin were eye pain, intraocular pressure increased, eye irritation (burning upon instillation) and eye pruritus occurring in less than 1% of patients.

Tabulated list of adverse reactions.

The following adverse reactions have been reported with Tobramycin during clinical trials or during post marketing experience and are classified according to the subsequent convention: very

common (≥ 1/10), common (≥ 1/100 to < 1/10), uncommon (≥ 1/1,000 to < 1/100), rare (≥ 1/10,000 to < 1/1,000) and very rare (< 1/10,000), and not known (cannot be estimated from the available data). Within each frequency-grouping, adverse reactions are presented in order of decreasing seriousness.

System organ classification	**Frequency**	**Adverse reaction**
Immune system disorders	Not known	anaphylactic reaction, hypersensitivity
Endocrine disorders	Not known	Cushing's syndrome, adrenal suppression (see section 4.4)
Nervous system disorders	Uncommon Not known	headache dizziness
Eye disorders	Uncommon Rare Not known	eye pain, eye pruritus, ocular discomfort, ocular hypertension, conjunctival oedema, increased intraocular pressure, eye irritation keratitis, eye allergy, vision blurred (see also section 4.4), dry eye, ocular hyperaemia eyelid oedema, erythema of the eyelid, mydriasis, lacrimation increased
Respiratory, thoracic, and mediastinal disorders	Uncommon	Rhinorrhoea, laryngospasm
Gastrointestinal disorders	Rare Not known	dysgeusia nausea, abdominal discomfort
Skin and subcutaneous tissue disorders	Not known	erythema multiforme, rash, swelling face, pruritus

Description of selected adverse reactions

The following adverse reactions have been observed following use with tobramycin ophthalmic solution:

System organ classification	**Frequency**	**Adverse reaction**
Eye disorders	Common Uncommon	ocular hyperaemia,* eye pain* eye pruritus,* ocular discomfort,* eye allergy, eyelid oedema,* conjunctivitis,* glare, increased lacrimation,* keratitis*

These adverse reactions were also observed with Tobramycin during post marketing.

Prolonged use of topical ophthalmic corticosteroids may result in increased intraocular pressure with damage to the optic nerve, reduced visual acuity and visual field defects, posterior subcapsular cataract formation and delayed wound healing.

Due to the corticosteroid component, in diseases causing thinning of the cornea or sclera there is a higher risk for perforation especially after long treatments (see section 4.4).

The development of secondary infection has occurred after the use of combinations containing corticosteroids and antimicrobials. Fungal infections of the cornea are particularly prone to develop coincidentally with long term applications of steroids.

Serious adverse reactions including neurotoxicity, ototoxicity and nephrotoxicity have occurred in patients receiving systemic tobramycin therapy (see section 4.4).

Sensitivity to topically administered aminoglycosides may occur in some patients (see section 4.4).

4.9 Overdose

There is no experience with overdosage with Artemether. There is no specific antidote known for the Artemisinin derivatives. However, experimental toxicological results obtained with large doses of Artemisinin on the cardiovascular system and the CNS should be considered. Overdosage could bring on cardiac irregularities. An ECG should be taken before initiating treatment in cardiac patients.

5. PHARMACOLOGICAL PROPERTIES

5.1 Pharmacodynamic properties

Pharmacotherapeutic group: Anti-inflammatory agents and anti-infectives in combination, corticosteroids and anti-infectives in combination

ATC code:S01AA12

Tobramycin:

Tobramycin is a potent, broad-spectrum, rapidly bactericidal aminoglycoside antibiotic. It exerts its primary effect on bacterial cells by inhibiting polypeptide assembly and synthesis on the ribosome. Tobramycin in this combination provides antibacterial protection against susceptible bacteria.

The following MIC breakpoints, separating susceptible from intermediate susceptible organisms, and intermediate susceptible from resistant organisms, are suggested: S ($\leq$ 4 µg/ml, R ($\geq$ 8 µg/ml). The prevalence of resistance may vary geographically and with time for selected species and local information on resistance is desirable, particularly when treating severe infections. As necessary, expert advice should be sought when the local prevalence of resistance is such that the utility of the agent in at least some types of infections is questionable. The following information gives only an approximate guidance on probabilities whether bacteria will be susceptible to tobramycin in Tobramycin.

The breakpoint definitions classifying isolates as susceptible or resistant are useful in predicting clinical efficacy of antibiotics that are administered systemically. However, when the antibiotic is administered in very high concentrations topically directly on the site of infection, these breakpoint definitions may not be applicable. Most isolates that would be classified as resistant by systemic breakpoints are indeed successfully treated topically.

In vitro studies have shown tobramycin to be active against most strains of common ocular pathogens and common skin flora bacteria as listed in the Table below:

Categories	**Frequency of Acquired Resistance in Europe**
SENSITIVE SPECIES	0-3%
Aerobic Gram-Positive Microorganisms	0-3%
Corynebacterium species	0-28%
Staphylococcus aureus Methicillin -S [a]	0-40%

Staphylococcus epidermidis Methicillin -S [a] Other Coagulase-negative Staphylococci	
Aerobic Gram-Negative Microorganisms *Acinetobacter* species **Citrobacter *species***	0% 0%
Escherichia coli *Enterobacter* species *Haemophilus influenzae*	0% 0% 0%
Klebsiella species *Moraxella* species *Proteus* species ***Pseudomonas aeruginosa***	0% 0% 0% 0%
MODERATELY SUSCEPTIBLE SPECIES **(*in vitro,* intermediate susceptibility)** ***Aerobic Gram-Negative Microorganisms*** *Serratia marcescens*	
INHERENTLY RESISTANT SPECIES ***Aerobic Gram-Positive Microorganisms*** Enterococcus species *Staphylococcus aureus* Methicillin –R [a] *Staphylococcus epidermidis* Methicillin –R [a] *Streptococcus pneumoniae* *Streptococcus* species	50 – 70% 30 – 40%
Aerobic Gram-negative microorganisms *Burkholderia cepacia* *Stenotrophomonas maltophilia*	
Anaerobic microorganisms Strict anaerobic bacteria	
Others *Chlamydia* species *Mycoplasma* species *Rickettsia* species	

[a] Methicillin-susceptible (S), Methicillin-resistant (R). The beta-lactam (i.e., methicillin; penicillin) resistance phenotype is unrelated to the aminoglycoside resistance phenotype and both are unrelated to the virulence phenotypes. Some methicillin-resistant (R) *S. aureus* strains (MRSA) are susceptible to tobramycin (MIC: S $\leq$ 4); conversely some strains of methicillin–susceptible (S) *S. aureus* (MSSA) are resistant to tobramycin (MIC: S $\geq$ 8)

The frequency of methicillin resistance (R) may be up to 50% of all staphylococci in some European countries

Paediatric population

The safety and efficacy of Tobramycin in children have been established by broad clinical experience, but only limited data are available. In a clinical study of Tobramycin suspension for the treatment of bacterial conjunctivitis, 29 paediatric patients, ranging in age from 1 to 17 years, were treated with 1 or 2 drops of Tobramycin every 4 or 6 hours for 5 or 7 days. In this study, differences in the safety profile between adult and paediatric patients were not observed.

Other information

Cross-resistance between aminoglycosides (e.g., gentamicin and tobramycin) is due to the specificity of the enzyme modifications, Adenyltransferase (ANT) and Acetyltransferase (ACC). However, cross-resistance varies between the aminoglycoside antibiotics due to the differing specificity of the various modifying enzymes. The most common mechanism of acquired resistance to aminoglycosides is antibiotic inactivation by plasmid and transposon-encoded modifying enzymes.

5.2 Pharmacokinetic properties

Tobramycin:

Animal studies have shown that tobramycin is absorbed into the cornea following ocular administration. Following systemic administration to patients with normal renal function, a plasma half-life of approximately 2 hours has been observed. Tobramycin is eliminated almost exclusively by glomerular filtration with little if any biotransformation. Plasma concentrations of tobramycin following the 2-day topical ocular regimen of Tobramycin were below the limit of quantification in most subjects or low (≤ 0.25 microgram/ml).

5.3 Preclinical safety data

Non-clinical data revealed no special hazard for humans from topical ocular exposure to tobramycin. Tobramycin has not been shown to induce teratogenicity in rats or rabbits.

6. PHARMACEUTICAL PARTICULARS

6.1 List of Excipients

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6.2 Incompatibilities

None

6.3 Shelf life

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6.4 Special precautions for storage

Store in the original package.

Store below 30°C. Keep medicine away from direct sunlight

Keep all medicine out of the reach of children.

6.5 Nature and contents of container

10ml dropper bottle placed in an inner carton with insert

6.6 Special precautions for disposal

No special requirements.
Any unused product or waste material should be disposed of in accordance with local requirements.

7. SUPPLIER AND MANUFACTURER

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