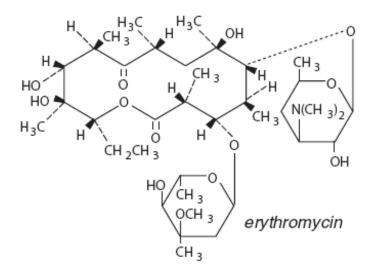
ERYTHROMYCIN- erythromycin ointment Lake Erie Medical DBA Quality Care Products LLC

Erythromycin Ophthalmic Ointment, USP 0.5% (Sterile)

Rx only

DESCRIPTION

Erythromycin Ophthalmic Ointment, USP belongs to the macrolide group of antibiotics. The sterile ophthalmic ointment flows freely over the conjunctiva. Erythromycin base, as crystals or powder, is slightly soluble in water, moderately soluble in ether, and readily soluble in alcohol or chloroform. Erythromycin is an antibiotic produced from a strain of *Streptomyces erythraeus*. It is basic and readily forms a salt when combined with an acid. It has the following structural formula:



 $C_{37}H_{67}NO_{13}$

Mol. Wt. 733.94

Chemical Name: (3R*, 4S*, 5S*, 6R*, 7R*, 9R*, 11R*, 12R*, 13S*, 14R*)-4-[(2,6-Dideoxy-3-C-methyl-3- θ -methyl- θ -meth

Each Gram Contains: ACTIVE: Erythromycin, USP 5 mg (0.5%); INACTIVES: Mineral Oil, White Petrolatum.

CLINICAL PHARMACOLOGY

Microbiology

Erythromycin inhibits protein synthesis without affecting nucleic acid synthesis. Erythromycin is usually active against the following organisms *in vitro* and in clinical infections: *Streptococcus pyogenes* (group A β-hemolytic), Alpha-hemolytic streptococci (viridans group); *Staphylococcus aureus*, including penicillinase-producing strains (methicillin-resistant staphylococci are uniformly resistant to erythromycin); *Streptococcus pneumoniae*; *Mycoplasma pneumoniae* (Eaton Agent, PPLO); *Haemophilus influenzae* (not all strains of this organism are susceptible at the erythromycin concentrations ordinarily achieved); *Treponema pallidum*; *Corynebacterium diphtheriae*; *Neisseria gonorrhoeae*; *Chlamydia trachomatis*.

INDICATIONS AND USAGE

For the treatment of superficial ocular infections involving the conjunctiva and/or cornea caused by organisms susceptible to erythromycin.

For prophylaxis of ophthalmia neonatorum due to *N. gonorrhoeae* or *C. trachomatis*.

The effectiveness of erythromycin in the prevention of ophthalmia caused by penicillinase-producing *N. gonorrhoeae* is not established.

For infants born to mothers with clinically apparent gonorrhea, intravenous or intramuscular injections of aqueous crystalline penicillin G should be given; a single dose of 50,000 units for term infants or 20,000 units for infants of low birth weight. Topical prophylaxis alone is inadequate for these infants.

CONTRAINDICATIONS

This drug is contraindicated in patients with a history of hypersensitivity to erythromycin.

PRECAUTIONS

General

The use of antimicrobial agents may be associated with the overgrowth of non-susceptible organisms including fungi; in such a case, antibiotic administration should be stopped and appropriate measures taken.

Information for Patients:

Avoid contaminating the tip of container with material from the eye, fingers or other source.

Carcinogenesis, Mutagenesis, Impairment of Fertility:

Two year oral studies conducted in rats with erythromycin did not provide evidence of tumorigenicity. Mutagenicity studies have not been conducted. No evidence of impaired fertility or harm to the fetus that appeared related to erythromycin was reported in these studies.

Pregnancy:

Pregnancy Category B.

Reproduction studies have been performed in rats, mice, and rabbits using erythromycin and its various salts and esters, at doses that were several multiples of the usual human dose. There are, however, no adequate and well-controlled studies in pregnant women. Because animal reproductive studies are not always predictive of human response, the erythromycins should be used during pregnancy only if clearly needed.

Nursing Mothers:

Caution should be exercised when erythromycin is administered to a nursing woman.

Pediatric Use:

See INDICATIONS AND USAGE and DOSAGE AND ADMINISTRATION.

ADVERSE REACTIONS

The most frequently reported adverse reactions are minor ocular irritations, redness and hypersensitivity reactions.

To report SUSPECTED ADVERSE REACTIONS, contact Bausch + Lomb, a division of Valeant Pharmaceuticals North America LLC at 1-800-321-4576 or FDA at 1-800-FDA-1088 or www.fda.gov/medwatch.

DOSAGE AND ADMINISTRATION

In the treatment of superficial ocular infections, erythromycin ophthalmic ointment approximately 1 cm in length should be applied directly to the infected eye(s) up to six times daily, depending on the severity of the infection.

For prophylaxis of neonatal gonococcal or chlamydial ophthalmia, a ribbon of ointment approximately 1 cm in length should be instilled into each lower conjunctival sac. The ointment should not be flushed from the eye following instillation. A new tube should be used for each infant.

HOW SUPPLIED

Erythromycin Ophthalmic Ointment, USP 0.5% is available in the following sizes:

1/8 oz. (3.5 g) tamper-resistant tube - NDC 55700-478-35

DO NOT USE IF CAP AND NECKRING ARE NOT INTACT.

DO NOT USE IF BOTTOM RIDGE OF TUBE CAP IS EXPOSED.

Storage:

Store between 15°-25°C (59°-77°F).

KEEP OUT OF REACH OF CHILDREN.

Revised: July 2016

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9043404 (Folded) 9043504 (Flat)



Packaged and Distributed by Zuality Care Products, LLC

1-800-337-8603



Holland, OH 43528

ERYTHROMYCIN OPHTHALMIC **OINTMENT USP, 0.5% (STERILE)**

GTIN: 00355700478353

NDC: 55700-0478-35

Serial:

1/8 oz (3.5g)

Lot:

EXP: //

Rx only

Each gram contains ACTIVE: Erythromycin, USP 5 mg (0.5%)

Mfr: Bausch & Lomb Incorporated, Tampa, FL 33637

Warning: Keep out of children's reach Store at controlled temperature 69-77 degrees F Consult with a physician or see insert for dosage



NDC: 55700-0478-35

ERYTHROMYCIN

erythromycin ointment

Product	Information
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Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:55700-478(NDC:24208-910)
Route of Administration	ОРНТНАЬМІС		

Active Ingredient/Active Moiety

Active ingrediction violety				
Ingredient Name	Basis of Strength	Strength		
ERYTHROMYCIN (UNII: 63937KV33D) (ERYTHROMYCIN - UNII:63937KV33D)	ERYTHROMYCIN	5 mg in 1 g		

Inactive Ingredients

inactive ingredients			
Ingredient Name	Strength		
PETROLATUM (UNII: 4T6H12BN9U)			
MINERAL O IL (UNII: T5L8T28FGP)			

Packaging

# Item Code	Package Description	Marketing Start Date	Marketing End Date
1 NDC:55700-478-35	1 in 1 CARTON	0 1/13/20 17	
1	3.5 g in 1 TUBE; Type 0: Not a Combination Product		

Marketing Information

Mari Neuring Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA064067	07/29/1994	

Labeler - Lake Erie Medical DBA Quality Care Products LLC (831276758)

Establishment			
Name	Address	ID/FEI	Business Operations
Lake Erie Medical DBA Quality Care Products LLC		831276758	relabel(55700-478)

Revised: 3/2019

Lake Erie Medical DBA Quality Care Products LLC