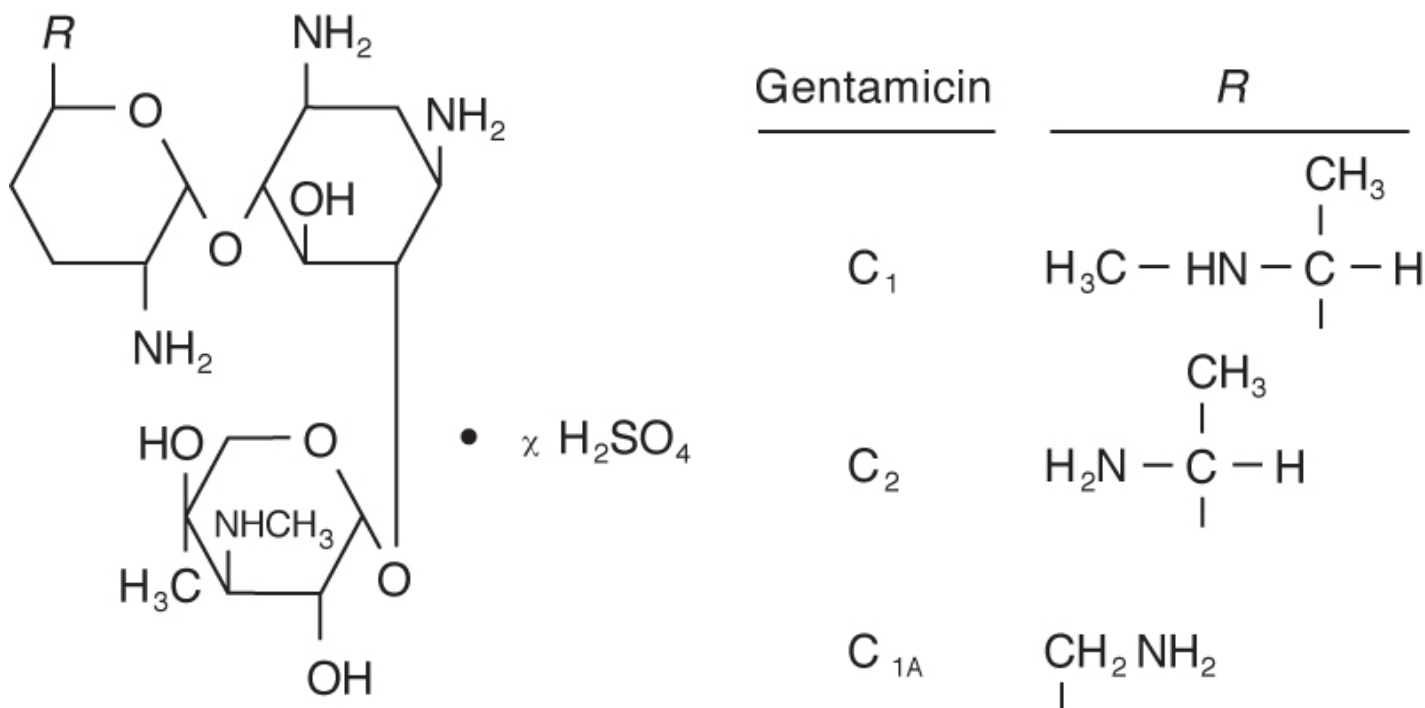


GENTAMICIN SULFATE- gentamicin sulfate solution
RPK Pharmaceuticals, Inc.

Gentamicin Sulfate
Ophthalmic Solution USP
Sterile
Rx Only

DESCRIPTION

Gentamicin Sulfate is a water-soluble antibiotic of the aminoglycoside group. Gentamicin Sulfate Ophthalmic Solution is a sterile, aqueous solution buffered to approximately pH 7 for ophthalmic use. Gentamicin is obtained from cultures of *Micromonospora purpurea*. It is a mixture of the sulfate salts of gentamicin C₁, C₂, and C_{1A}. All three components appear to have similar antimicrobial activities. Gentamicin sulfate occurs as white powder and is soluble in water and insoluble in alcohol. The structure is as follows:



Established name: Gentamicin Sulfate

Chemical name: 0-3-Deoxy-4-C-methyl-3-(methylamino)-β-L-arabinopyranosyl-(1 → 6)-0-[2,6-diamino-2,3,4,6-tetrahydroxy-α-D-erythro-hexopyranosyl-(1 → 4)]-2-deoxy-D-streptamine.

Ingredients: Each mL contains: Active: Gentamicin Sulfate USP (equivalent to 3 mg gentamicin).

Preservative: Benzalkonium Chloride 0.1 mg (0.01%). **Inactives:** Sodium Chloride, Dried Sodium Phosphate, Tyloxapol, Sodium Hydroxide and/or Hydrochloric Acid (to adjust pH) and Purified Water.

CLINICAL PHARMACOLOGY

Microbiology: Gentamicin sulfate is active *in vitro* against many strains of the following microorganisms:

Staphylococcus aureus, Staphylococcus epidermidis, Streptococcus pyogenes, Streptococcus pneumoniae, Enterobacter aerogenes, Escherichia coli, Haemophilus influenzae, Klebsiella pneumoniae, Neisseria gonorrhoeae, Pseudomonas aeruginosa, and Serratia marcescens.

INDICATIONS AND USAGE

Gentamicin Sulfate Ophthalmic Solution is indicated in the topical treatment of ocular bacterial infections, including conjunctivitis, keratitis, keratoconjunctivitis, corneal ulcers, blepharitis, blepharoconjunctivitis, acute meibomianitis, and dacryocystitis caused by susceptible strains of the following microorganisms: *Staphylococcus aureus, Staphylococcus epidermidis, Streptococcus pyogenes, Streptococcus pneumoniae, Enterobacter aerogenes, Escherichia coli, Haemophilus influenzae, Klebsiella pneumoniae, Neisseria gonorrhoeae, Pseudomonas aeruginosa, and Serratia marcescens.*

CONTRAINDICATIONS

Gentamicin Sulfate Ophthalmic Solution is contraindicated in patients with known hypersensitivity to any of the components.

WARNINGS

NOT FOR INJECTION INTO THE EYE. Gentamicin Sulfate Ophthalmic Solution is not for injection. It should never be injected subconjunctivally, nor should it be directly introduced into the anterior chamber of the eye.

PRECAUTIONS

General

Prolonged use of topical antibiotics may give rise to overgrowth of non-susceptible organisms including fungi. Bacterial resistance to gentamicin may also develop. If purulent discharge, inflammation or pain becomes aggravated, the patient should discontinue use of the medication and consult a physician. If irritation or hypersensitivity to any component of the drug develops, the patient should discontinue use of this preparation, and appropriate therapy should be instituted.

Information for Patients

To avoid contamination, do not touch tip of container to the eye, eyelid, or any surface.

Carcinogenesis, Mutagenesis, Impairment of Fertility

There are no published carcinogenicity or impairment of fertility studies on gentamicin. Aminoglycoside antibiotics have been found to be non-mutagenic.

Pregnancy

Pregnancy Category C. Gentamicin has been shown to depress body weights, kidney weights, and median glomerular counts in newborn rats when administered systemically to pregnant rats in daily doses approximately 500 times the maximum recommended ophthalmic human dose. There are no adequate and well-controlled studies in pregnant women. Gentamicin should be used during pregnancy only if the potential benefit justifies the potential risk to the fetus.

Pediatric Use

Safety and effectiveness in neonates have not been established.

ADVERSE REACTIONS

Bacterial and fungal corneal ulcers have developed during treatment with gentamicin ophthalmic preparations. The most frequently reported adverse reactions are ocular burning and irritation upon drug instillation, non-specific conjunctivitis, conjunctival epithelial defects, and conjunctival hyperemia. Other adverse reactions which have occurred rarely are allergic reactions, thrombocytopenic purpura, and hallucinations.

To report SUSPECTED ADVERSE REACTIONS, contact Sandoz Inc. at 1-800-525-8747 or FDA at 1-800-FDA-1088 or www.fda.gov/medwatch.

DOSAGE AND ADMINISTRATION

Instill one or two drops into the affected eye(s) every four hours. In severe infections, dosage may be increased to as much as two drops once every hour.

HOW SUPPLIED

Product: 53002-9291

NDC: 53002-9291-1 5 mL in a BOTTLE, DROPPER

Manufactured by

Alcon Laboratories, Inc.

Fort Worth, Texas 76134 for

Sandoz Inc.

Princeton, NJ 08540

Rev. June 2020

300036475-0620

Gentamicin Ophthalmic Solution 0.3%, USP

The image shows several labels for Gentamicin Ophthalmic Solution 0.3%, USP. The main label features the following text:

- 5 mL Bottle RxC# 191880431000 ORDER# K33-71**
- GENTAMICIN 0.3% OPHTHAL SOLN**
- SANDOZ** Generic for GARAMYCIN
- INSTILL 1-2 DROPS INTO AFFECTED EYE(S) 4 TIMES A DAY OR AS DIRECTED.**
- Rx only**
- LOT# 61314-633-05**
- EXP: 10-31-2021**
- ORDER# K33-71**
- 5 mL GENTAMICIN 0.3% OPHTHAL SOLN**
- 1-2 GTTS AFF EYES QID**
- LOT# 19123-014 EXP 10-31-2021**
- RxC# 191880431000 FDA-9291**
- 5 mL GENTAMICIN 0.3% OPHTHAL SOLN**
- BILLING NDC# 51014-0633-05**
- RxC# 191880431000**
- 5 mL GENTAMICIN 0.3% OPHTHAL SOLN**
- BILLING NDC# 51014-0633-05**
- RxC# 191880431000**
- 5 mL GENTAMICIN 0.3% OPHTHAL SOLN**
- BILLING NDC# 51014-0633-05**
- RxC# 191880431000**
- 5 mL GENTAMICIN 0.3% OPHTHAL SOLN**
- GENTAMICIN OPHTHALMIC SOLUTION 0.3%, USP**
- 5 mL Bottle**
- DISCARD BY 10-31-2021**
- NDC# 53002-9291-1 RxC# 191880431000**
- FEDERAL LAW PROHIBITS TRANSFER OF THIS MEDICATION ANY PERSON OTHER THAN THE PERSON FOR WHOM IT WAS PRESCRIBED. KEEP THIS & ALL MEDICATIONS OUT OF THE REACH OF CHILDREN. CALL YOUR DOCTOR FOR MEDICAL ADVICE ABOUT SIDE EFFECTS. YOU MAY REPORT SIDE EFFECTS TO FDA AT 1-800-FDA-8088.**
- Clinic Name Here**
- PRESCRIBER NAME DATE**
- PATIENT NAME**

GENTAMICIN SULFATE

gentamicin sulfate solution

Product Information

Product Type

HUMAN PRESCRIPTION DRUG

Item Code (Source)

NDC:53002-9291(NDC:61314-633)

Route of Administration

OPHTHALMIC

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
GENTAMICIN SULFATE (UNII: 8X7386QRLV) (GENTAMICIN - UNII:T6Z9V48IKG)	GENTAMICIN	3 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
SODIUM PHOSPHATE (UNII: SE337SVY37)	
TYLOXAPOL (UNII: Y27PUL9H56)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
HYDROCHLORIC ACID (UNII: QTT17582CB)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:53002-9291-1	5 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	01/01/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA062196	04/05/1996	

Labeler - RPK Pharmaceuticals, Inc. (147096275)**Establishment**

Name	Address	ID/FEI	Business Operations
RPK Pharmaceuticals, Inc.		147096275	RELABEL(53002-9291) , REPACK(53002-9291)

Revised: 12/2020

RPK Pharmaceuticals, Inc.