# NEOMYCIN AND POLYMYXIN B SULFATES AND DEXAMETHASONE- neomycin sulfate, polymyxin b sulfate, and dexamethasone ointment Proficient Rx LP

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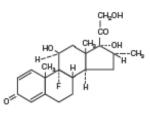
Neomycin and Polymyxin B Sulfates and Dexamethasone Ophthalmic Ointment USP

(Sterile)

**Rx only** 

#### **DESCRIPTION:**

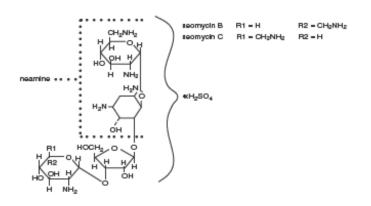
Neomycin and Polymyxin B Sulfates and Dexamethasone Ophthalmic Ointment USP is a multiple dose anti-infective steroid combination in a sterile ointment for topical application. The active ingredient dexamethasone, is represented by the following structural formula:



C<sub>22</sub>H<sub>29</sub>FO<sub>5</sub> Mol. Wt. 392.47

Chemical Name: Pregna-1,4-diene-3,20-dione, 9-fluoro-11,17,21-trihydroxy-16-methyl-, (11β,16α)-.

Neomycin Sulfate is the sulfate salt of neomycin B and C which are produced by the growth of *Streptomyces fradiae* Waksman (Fam. Streptomycetaceae). It has a potency equivalent to not less than 600 micrograms of neomycin base per milligram, calculated on an anhydrous basis. The structural formulae are:



Polymyxin B Sulfate is the sulfate salt of polymyxin  $B_1$  and  $B_2$  which are produced by the growth of *Bacillus polymyxa* (Prazmowski) Migula (Fam. Bacillaceae). It has a potency of not less than 6,000 polymyxin B units per milligram, calculated on an anhydrous basis.

The structural formulae are:

**Each Gram Contains:** ACTIVES: Neomycin Sulfate (equivalent to 3.5 mg Neomycin), Polymyxin B Sulfate, equal to 10,000 polymyxin B units; Dexamethasone 1 mg (0.1%). INACTIVES: White Petrolatum, Lanolin, Mineral Oil.

PRESERVATIVES ADDED: Methylparaben 0.05%, Propylparaben 0.01%.

#### **CLINICAL PHARMACOLOGY:**

Corticoids suppress the inflammatory response to a variety of agents and they probably delay or slow the healing. Since corticoids may inhibit the body's defense mechanism against infections, a concomitant antimicrobial drug may be used when this inhibition is considered to be clinically significant in a particular case.

When a decision to administer both a corticoid and an antimicrobial is made, the administration of such drugs in combination has the advantage of greater patient compliance and convenience, with the added assurance that the appropriate dosage of both drugs is administered, plus assured compatibility of ingredients when both types of drugs are in the same formulation and, particularly, that the correct volume of drug is delivered and retained.

The relative potency of corticosteroids depends on the molecular structure, concentration, and release from the vehicle.

#### **INDICATIONS AND USAGE:**

For steroid-responsive inflammatory ocular conditions for which a corticosteroid is indicated and where bacterial infection or a risk of bacterial ocular infection exists.

Ocular steroids are indicated in inflammatory conditions of the palpebral and bulbar conjunctiva, cornea, and anterior segment of the globe where the inherent risk of steroid use in certain cases of infective conjunctivitides is accepted to obtain a diminution in edema and inflammation. They are also indicated in chronic anterior uveitis and corneal injury from chemical radiation or thermal burns; or penetration of foreign bodies.

The use of a combination drug with an anti-infective component is indicated where the risk of infection is high or where there is an expectation that potentially dangerous numbers of bacteria will be present in the eye.

The particular anti-infective drugs in this product is active against the following common bacterial eye pathogens: *Staphylococcus aureus*, *Escherichia coli*, *Haemophilus influenzae*, *Klebsiella/Enterobacter species*, *Neisseria* species, *Pseudomonas aeruginosa*.

This product does not provide adequate coverage against Serratia marcescens, and Streptococci, including *Streptococcus pneumoniae*.

#### **CONTRAINDICATIONS:**

Epithelial herpes simplex keratitis (dendritic keratitis), vaccinia, varicella and many other viral diseases of the cornea and conjunctiva. Mycobacterial infection of the eye. Fungal diseases of ocular structures.

Hypersensitivity to a component of the medication. (Hypersensitivity to the antibiotic component occurs at a higher rate than for other components.)

#### WARNINGS:

NOT FOR INJECTION. Do not touch tube tip to any surface, as this may contaminate the contents. Prolonged use may result in glaucoma, with damage to the optic nerve, defects in visual acuity and fields of vision, and posterior subcapsular cataract formation. Prolonged use may suppress the host response and thus increase the hazard of secondary ocular infections. In those diseases causing thinning of the cornea or sclera, perforations have been known to occur with the use of topical steroids. In acute purulent conditions of the eye, steroids may mask infection or enhance existing infection. If these products are used for 10 days or longer, intraocular pressure should be routinely monitored even though it may be difficult in children and uncooperative patients.

Products containing neomycin sulfate may cause cutaneous sensitization.

Employment of steroid medication in the treatment of herpes simplex requires great caution.

## **PRECAUTIONS:**

#### General

The initial prescription and renewal of the medication order beyond 8 grams should be made by a physician only after examination of the patient with the aid of magnification, such as slit lamp biomicroscopy and, where appropriate, fluorescein staining.

The possibility of persistent fungal infections of the cornea should be considered after prolonged steroid dosing.

#### **Information for Patients:**

Do not touch tube tip to any surface, as this may contaminate the contents.

#### **Pregnancy:**

**Pregnancy Category C.** Dexamethasone has been shown to be teratogenic in mice and rabbits following topical ophthalmic application in multiples of the therapeutic dose.

In the mouse, corticosteroids produce fetal resorptions and a specific abnormality, cleft palate. In the rabbit, corticosteroids have produced fetal resorptions and multiple abnormalities involving the head, ears, limbs, palate, etc.

There are no adequate or well-controlled studies in pregnant women. Neomycin, Polymyxin B Sulfates and Dexamethasone Ophthalmic Ointment should be used during pregnancy only if the potential benefit to the mother justifies the potential risk to the embryo or fetus. Infants born of mothers who have received substantial doses of corticosteroids during pregnancy should be observed carefully for signs of hypoadrenalism.

## Nursing Mothers:

Systemically administered corticosteroids appear in human milk and could suppress growth, interfere with endogenous corticosteroid production, or cause other untoward effects. It is not known whether topical administration of corticosteroids could result in sufficient systemic absorption to produce detectable quantities in human milk. Because many drugs are excreted in human milk, caution should be exercised when Neomycin, Polymyxin B Sulfates and Dexamethasone Ophthalmic Ointment, is administered to a nursing woman.

## **Pediatric Use:**

Safety and effectiveness in pediatric patients have not been established.

# **ADVERSE REACTIONS:**

Adverse reactions have occurred with steroid/anti-infective combination drugs which can be attributed to the steroid component, the anti-infective component, or the combination. Exact incidence figures are not available since no denominator of treated patients is available.

Reactions occurring most often from the presence of the anti-infective ingredients are allergic sensitizations. The reactions due to the steroid component are: elevation of intraocular pressure (IOP) with possible development of glaucoma, and infrequent optic nerve damage; posterior subcapsular cataract formation; and delayed wound healing.

**Secondary Infection:** The development of secondary infection has occurred after use of combinations containing steroids and antimicrobials. Fungal infections of the cornea are particularly prone to develop coincidentally with long-term applications of steroids. The possibility of fungal invasion must be considered in any persistent corneal ulceration where steroid treatment has been used.

Secondary bacterial ocular infection following suppression of host responses also occurs.

# DOSAGE AND ADMINISTRATION:

Apply a small amount into the conjunctival sac(s) up to three or four times daily, or may be used adjunctively with Neomycin and Polymyxin B Sulfates and Dexamethasone Ophthalmic Suspension at bedtime.

How to apply Neomycin and Polymyxin B Sulfates and Dexamethasone Ophthalmic Ointment:

1. Tilt your head back.

2. Place a finger on your cheek just under your eye and gently pull down until a "V" pocket is formed between your eyeball and your lower lid.

3. Place a small amount (about 1/2 inch) of Neomycin and Polymyxin B Sulfates and Dexamethasone Ophthalmic Ointment in the "V" pocket. Do not let the tip of the tube touch your eye.

4. Look downward before closing your eye.

Not more than 8 grams should be prescribed initially and the prescription should not be refilled without further evaluation as outlined in PRECAUTIONS above.

## HOW SUPPLIED:

Neomycin and Polymyxin B Sulfates and Dexamethasone Ophthalmic Ointment USP is supplied in an ophthalmic tube in the following size:

3.5 gram tube – Prod. No. 04034

## Storage:

Store between 15°-25° C (59°-77° F).

## KEEP OUT OF REACH OF CHILDREN.

# DO NOT USE IF CAP AND NECKRING ARE NOT INTACT.

## FOR OPHTHALMIC USE ONLY.

Revised November 2007

**Bausch & Lomb Incorporated** Tampa, FL 33637 ©Bausch & Lomb Incorporated

Relabeled by:

**Proficient Rx LP** Thousand Oaks, CA 91320

9114002 (Folded) 9114102 (Flat)

# **Principal Display Panel**

ProficientRy	NDC 63187-669-35	Lot #:00000 Exp. 00/00/00 SN# MASTER	
	RX Only	Neomycin / Poly B / Dexamethasone 3.5gm Ophthalmic Ointment Lot #:0000 SN# MASTER NDC 63187-669-35 Exo:00/00/00	
Neomycin / Poly B / Dexa			
3.5gm Ophthalmic Ointment		Neomycin / Poly B / Dexamethasone 3.5gm Ophthalmic Ointment	
Each gram contains: Neomycin Sulfate ( equivalent to 3.5mg Neomycin), Polymyxin B Sulfate, equal to		Lot #:00000 SN# MASTER NDC 63187-669-35 Exp:00/00/00	
10,000 Polymyxin B units, Dexa (0.1%)	amethasone 1 mg	Neomycin / Poly B / Dexamethasone 3.5gm Ophthalmic Ointment Lot #00000 SN# MASTER	
See package insert. FOR OPHTHALMIC USE ONLY.		NDC 63187-669-35 Exp:00/00/00	
Product ID: RN066935			
Mfr. By: Bausch & Lomb Incorporated Tampa, FL 33637	(eep medication out of the reach of children	Relabeled By: Proficient Rx LP	
Store between 15°-25° C (59°-77° F).	Thousand Oaks, CA 91320		

NDC 63187-669-35

Neomycin and Polymyxin B Sulfates and Dexamethasone Ophthalmic Ointment USP

(Sterile)

Rx only

[icon- eye] [icon- ointment] [icon- 3.5gm]

Net wt. 1/8 oz (3.5g)

# NEOMYCIN AND POLYMYXIN B SULFATES AND DEXAMETHASONE

neomycin sulfate, polymyxin b sulfate, and dexamethasone ointment

Product Information					
Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:63187-669(NDC:24208-795)		
Route of Administration	OPHTHALMIC				
Active Ingredient/Active Moiety					

	Ingredient Name	Basis of Stren	gth Strength
EOMYCIN SULFATE (UNII: 057Y626693) (NEOMYCIN - UNII:116QD7X297) NEOMYCIN			3.5 mg in 1 g
POLYMYXIN B SULFATE (UNII: 19371312D4) (POLYMYXIN B - UNII:J2VZ07J96K) POLYMYXIN B			10000 [USP'U] in 1 g
DEXAMETHASONE (U	L) DEXAMETHASO	NE 1 mg in 1 g	
Inactive Ingredie	nts		
Ingredient Name			Strength
LANOLIN (UNII: 7EV65	5EAW6H)		
METHYLPARABEN (U			
MINERAL OIL (UNII: T			
PROPYLPARABEN (UI	NII: Z8IX2SC1OH)		
	,		
PROPYLPARABEN (UI	,		
PROPYLPARABEN (UI	,		
PROPYLPARABEN (UI PETROLATUM (UNII: 4	,	Marketing Start Date	e Marketing End Date
PROPYLPARABEN (UI PETROLATUM (UNII: 4 Packaging	4T6H12BN9U) Package Description	Marketing Start Date	e Marketing End Date
PROPYLPARABEN (UI PETROLATUM (UNII: 4 Packaging # Item Code	4T6H12BN9U) Package Description		e Marketing End Date
PROPYLPARABEN (UI PETROLATUM (UNII: 4 Packaging # Item Code 1 NDC:63187-669-35	4T6H12BN9U) Package Description 1 in 1 CARTON		e Marketing End Date
PROPYLPARABEN (UI PETROLATUM (UNII: 4 Packaging # Item Code 1 NDC:63187-669-35	4T6H12BN9U) Package Description 1 in 1 CARTON		e Marketing End Date
PROPYLPARABEN (UI PETROLATUM (UNII: 4 PETROLATUM (UNII: 4 I tem Code NDC:63187-669-35 I	TGH12BN9U) Package Description 1 in 1 CARTON 3.5 g in 1 TUBE; Type 0: Not a Combination Product		e Marketing End Date
PROPYLPARABEN (UI PETROLATUM (UNII: 4 Packaging # Item Code 1 NDC:63187-669-35	Package Description 1 in 1 CARTON 3.5 g in 1 TUBE; Type 0: Not a Combination Product prmation		
PROPYLPARABEN (U) PETROLATUM (UNII: 4 Packaging # Item Code 1 NDC:63187-669-35 1 NDC:63187-669-35	Package Description 1 in 1 CARTON 3.5 g in 1 TUBE; Type 0: Not a Combination Product prmation	12/0 1/20 18	

Labeler - Proficient Rx LP (079196022)

# Establishment

Name	Address	ID/FEI	<b>Business Operations</b>
Proficient Rx LP		079502574	RELABEL(63187-669)

Revised: 12/2018

Proficient Rx LP