

**TOBRAMYCIN AND DEXAMETHASONE- tobramycin and dexamethasone suspension/ drops**  
**Amneal Pharmaceuticals NY LLC**

-----  
**Tobramycin and Dexamethasone Ophthalmic Suspension, USP**  
**Sterile**

**DESCRIPTION**

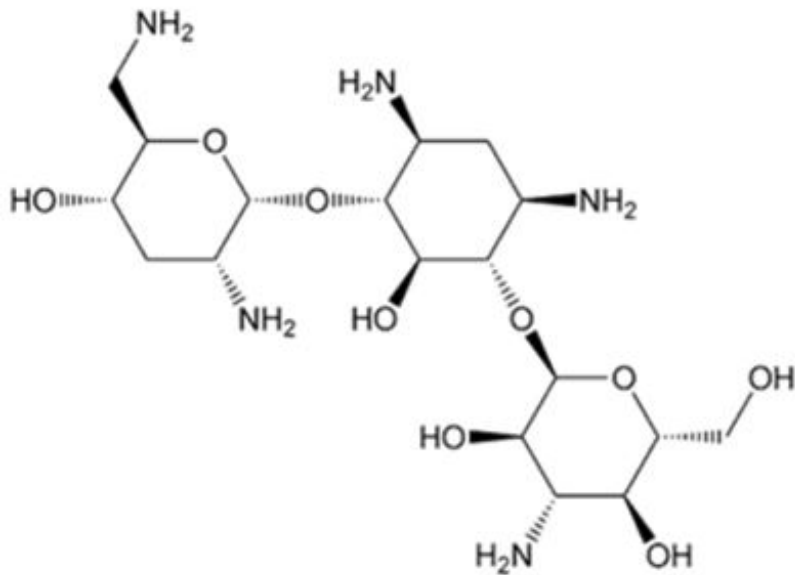
Tobramycin and dexamethasone ophthalmic suspension, USP is a sterile, multiple-dose antibiotic and steroid combination for topical ophthalmic use.

The chemical structures for tobramycin and dexamethasone are presented below:

**Tobramycin, USP**

Tobramycin, USP is a white to off-white, hygroscopic powder. It is freely soluble in water, very slightly soluble in alcohol, practically insoluble in chloroform and in ether.

Chemical Structure:



Molecular Formula: C<sub>18</sub>H<sub>37</sub>N<sub>5</sub>O<sub>9</sub>

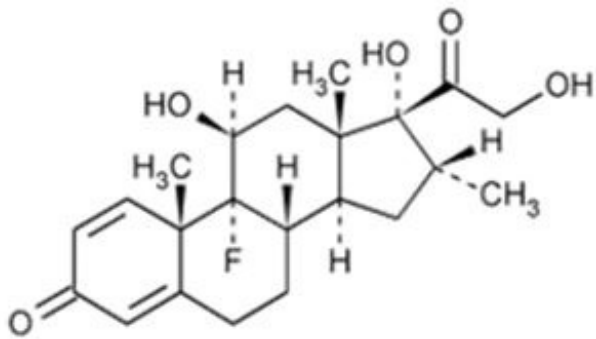
Molecular Weight: 467.51

Chemical Name: *O*-3-Amino-3-deoxy- $\alpha$ -D-glucopyranosyl-(1 $\rightarrow$ 4)-O-[2,6-diamino-2,3,6-trideoxy- $\alpha$ -D-ribo-hexopyranosyl (1 $\rightarrow$ 6)]-2-deoxy-L-streptamine

**Dexamethasone, USP**

Dexamethasone, USP (Micronised) is a white or practically white crystalline powder.

Chemical Structure:



Molecular Formula:  $C_{22}H_{29}FO_5$

Molecular Weight: 392.46

Chemical Name: 9-Fluoro-11 $\beta$ ,17,21-trihydroxy-16 $\alpha$ -methylpregna-1,4-diene-3,20-dione

**Each mL of tobramycin and dexamethasone ophthalmic suspension, USP contains:**

**Active:** tobramycin, USP 0.3% (3 mg) and dexamethasone, USP 0.1% (1 mg).

**Preservative:** benzalkonium chloride 0.01%.

**Inactives:** tyloxapol, edetate disodium, sodium chloride, hydroxyethyl cellulose, sodium sulfate, sulfuric acid and/or sodium hydroxide (to adjust pH) and water for injection, USP.

## CLINICAL PHARMACOLOGY

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

The antibiotic component in the combination (tobramycin) is included to provide action against susceptible organisms. *In vitro* studies have demonstrated that tobramycin is active against susceptible strains of the following microorganisms:

*Staphylococci*, including *S. aureus* and *S. epidermidis* (coagulase-positive and coagulase-negative), including penicillin-resistant strains.

*Streptococci*, including some of the Group A-beta-hemolytic species, some nonhemolytic species, and some *Streptococcus pneumoniae*.

*Pseudomonas aeruginosa*, *Escherichia coli*, *Klebsiella pneumoniae*, *Enterobacter aerogenes*, *Proteus mirabilis*, *Morganella morganii*, most *Proteus vulgaris* strains, *Haemophilus influenzae* and *H. aegyptius*, *Moraxella lacunata*, *Acinetobacter calcoaceticus* and some *Neisseria* species.

No data are available on the extent of systemic absorption from tobramycin and dexamethasone ophthalmic suspension; however, it is known that some systemic absorption can occur with ocularly applied drugs.

## INDICATIONS AND USAGE

Tobramycin and dexamethasone ophthalmic suspension is indicated for steroid-responsive inflammatory ocular conditions for which a corticosteroid is indicated and where superficial bacterial ocular 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 infective conjunctivitis 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 superficial ocular 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 drug in this product is active against the following common bacterial eye pathogens:

*Staphylococci*, including *S. aureus* and *S. epidermidis* (coagulase-positive and coagulase-negative), including penicillin-resistant strains.

*Streptococci*, including some of the Group A-beta-hemolytic species, some nonhemolytic species, and some *Streptococcus pneumoniae*.

*Pseudomonas aeruginosa*, *Escherichia coli*, *Klebsiella pneumoniae*, *Enterobacter aerogenes*, *Proteus mirabilis*, *Morganella morganii*, most *Proteus vulgaris* strains, *Haemophilus influenzae* and *H. aegyptius*, *Moraxella lacunata*, *Acinetobacter calcoaceticus* and some *Neisseria* species.

## **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.

## **WARNINGS**

FOR TOPICAL OPHTHALMIC USE. **NOT FOR INJECTION INTO THE EYE.**

Sensitivity to topically applied aminoglycosides may occur in some patients. Severity of hypersensitivity reactions may vary from local effects to generalized reactions, such as erythema, itching, urticaria, skin rash, anaphylaxis, anaphylactoid reactions, or bullous reactions. If a sensitivity reaction does occur, discontinue use.

Prolonged use of steroids may result in glaucoma, with damage to the optic nerve, defects in visual acuity and fields of vision, and posterior subcapsular cataract formation. Intraocular pressure (IOP) should be routinely monitored even though it may be difficult in pediatric patients and uncooperative patients. Prolonged use may suppress the host response and thus increase the hazard of secondary ocular infections. In acute purulent conditions and parasitic infections of the eye, steroids may mask infection or enhance existing infection.

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

## **PRECAUTIONS**

### **General**

The possibility of fungal infections of the cornea should be considered after long-term steroid dosing. As with other antibiotic preparations, prolonged use may result in overgrowth of nonsusceptible organisms, including fungi. If superinfection occurs, appropriate therapy should be initiated. When multiple prescriptions are required, or whenever clinical judgement dictates, the patient should be examined with the aid of magnification, such as slit lamp biomicroscopy and, where appropriate, fluorescein staining.

Cross-sensitivity to other aminoglycoside antibiotics may occur; if hypersensitivity develops with this product, discontinue use and institute appropriate therapy.

### **Information for Patients**

Do not touch dropper tip to any surface, as this may contaminate the contents. Contact lenses should not be worn during the use of this product.

### **Carcinogenesis, Mutagenesis, Impairment of Fertility**

No studies have been conducted to evaluate the carcinogenic or mutagenic potential. No impairment of fertility was noted in studies of subcutaneous tobramycin in rats at doses of 50 and 100 mg/kg/day.

### **Pregnancy**

Corticosteroids have been found to be teratogenic in animal studies. Ocular administration of 0.1% dexamethasone resulted in 15.6% and 32.3% incidence of fetal anomalies in two groups of pregnant rabbits. Fetal growth retardation and increased mortality rates have been observed in rats with chronic dexamethasone therapy. Reproduction studies have been performed in rats and rabbits with tobramycin at doses up to 100 mg/kg/day parenterally and have revealed no evidence of impaired fertility or harm to the fetus.

There are no adequate and well-controlled studies in pregnant women. However, prolonged or repeated corticoid use during pregnancy has been associated with an increased risk of intra-uterine growth retardation. Tobramycin and dexamethasone ophthalmic suspension should be used during pregnancy only if the potential benefit justifies the potential risk to the 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 tobramycin and dexamethasone ophthalmic suspension is administered to a nursing woman.

## **Pediatric Use**

Safety and effectiveness in pediatric patients below the age of 2 years have not been established.

## **Geriatric Use**

No overall differences in safety or effectiveness have been observed between elderly and younger patients.

## **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.

The most frequent adverse reactions to topical ocular tobramycin (tobramycin ophthalmic solution, 0.3%) are hypersensitivity and localized ocular toxicity, including lid itching and swelling, and conjunctival erythema. These reactions occur in less than 4% of patients. The reactions due to the steroid component are: elevation of 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.

## **Postmarketing Experience**

Additional adverse reactions identified from postmarketing use include anaphylactic reaction, erythema multiforme.

The following additional adverse reactions have been reported with the individual components listed below:

Dexamethasone: Cushing's syndrome and adrenal suppression may occur after use of dexamethasone in excess of the listed dosing instructions in predisposed patients, including children and patients treated with CYP3A4 inhibitors.

Aminoglycosides: Neurotoxicity, ototoxicity, and nephrotoxicity have occurred in patients receiving systemic aminoglycoside therapy. Aminoglycosides may aggravate muscle weakness in patients with known or suspected neuromuscular disorders, such as myasthenia gravis or Parkinson's disease, because of their potential effect on neuromuscular function.

**To report SUSPECTED ADVERSE REACTIONS, contact Amneal Pharmaceuticals at 1-877-835-5472 or FDA at 1-800-FDA-1088 or [www.fda.gov/medwatch](http://www.fda.gov/medwatch).**

## **DOSAGE AND ADMINISTRATION**

One or two drops instilled into the conjunctival sac(s) every four to six hours. During the initial 24 to 48 hours, the dosage may be increased to one or two drops every two (2) hours. Frequency should be decreased gradually as warranted by improvement in clinical signs. Care should be taken not to discontinue therapy prematurely.

Not more than 20 mL should be prescribed initially and the prescription should not be refilled without further evaluation as outlined in **PRECAUTIONS** above.

## **HOW SUPPLIED**

Tobramycin and Dexamethasone Ophthalmic Suspension USP, **0.3% and 0.1%** is a sterile, white to off-white uniform suspension, essentially free from foreign matter, supplied in 2.5 mL 5 mL or 10 mL LDPE natural bottle with LDPE natural nozzle and HDPE white cap. Each mL contains tobramycin, USP 0.3% (3 mg) and dexamethasone, USP 0.1% (1 mg).

It is available as follows:

2.5 mL in 5 mL Container:	NDC 69238-1373-2
5 mL in 5 mL Container:	NDC 69238-1374-3
10 mL in 10 mL Container:	NDC 69238-1375-6

## **STORAGE**

Store at 8° to 27°C (46° to 80°F).

Store suspension upright and shake well before using.

After opening, tobramycin and dexamethasone ophthalmic suspension can be used until the expiration date on the bottle.

Manufactured by:

**Amneal Pharmaceuticals Pvt. Ltd.**

**Parenteral Unit**

Ahmedabad 382213, INDIA

Distributed by:

**Amneal Pharmaceuticals LLC**

Bridgewater, NJ 08807

Rev. 05-2021-05

## **PRINCIPAL DISPLAY PANEL**

**NDC 69238-1373-2**

**Tobramycin and Dexamethasone Ophthalmic Solution USP, 0.3%/0.1%**

**Rx Only**

**2.5 mL**

**Amneal Pharmaceuticals LLC**

**Unvarnished Area for, (6 x 14 mm)**

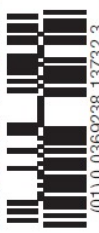
NDC 69238-1373-2  
**Tobramycin and Dexamethasone Ophthalmic Suspension, USP**  
**0.3%/0.1%**  
 For Topical Ophthalmic Use Only  
 Sterile  
 2.5 mL Rx only

**Each mL contains:**  
**Active:** Tobramycin, USP 0.3% (3 mg) and Dexamethasone, USP 0.1% (1 mg).  
**Preservative:** Benzalkonium chloride 0.01%. **Inactive:** Tyloxapol, edetate disodium, sodium chloride, hydroxyethyl cellulose, sodium sulfate, sulfuric acid and/or sodium hydroxide (to adjust pH) and water for injection, USP.

**PRECAUTION:** Do not touch dropper tip to any surface, as this may contaminate the suspension.  
**Store at 8° to 27°C (46° to 80°F). Store upright. SHAKE WELL BEFORE USING.**

**Usual Dosage:**  
 READ ENCLOSED INSERT.  
 Mfg. Lic. No. G/28/1539  
 Distributed by:  
**Anneal Pharmaceuticals LLC**  
 Bridgewater, NJ 08807  
 Rev. 04-2024-04 Made in INDIA

**Unvarnished Area for, Lot. & Exp. (10 x 14 mm)**




**Unvarnished Area for (38 mm x 35 mm)**  
 2D barcode along with GTIN, Lot, Exp. & Sr. No.

PMI-0569

Space for UPs No.

NDC 69238-1373-2  
**Tobramycin and Dexamethasone Ophthalmic Suspension, USP**  
**0.3%/0.1%**  
 For Topical Ophthalmic Use Only  
 Sterile  
 2.5 mL Rx only

**amneal**

**Each mL contains:**  
**Active:** Tobramycin, USP 0.3% (3 mg) and Dexamethasone, USP 0.1% (1 mg).  
**Preservative:** Benzalkonium chloride 0.01%.  
**Inactive:** Tyloxapol, edetate disodium, sodium chloride, hydroxyethyl cellulose, sodium sulfate, sulfuric acid and/or sodium hydroxide (to adjust pH) and water for injection, USP.  
**PRECAUTION:** Do not touch dropper tip to any surface, as this may contaminate the suspension.  
**Store at 8° to 27°C (46° to 80°F). Store upright.**  
**SHAKE WELL BEFORE USING.**  
**Usual Dosage:** READ ENCLOSED INSERT.


NDC 69238-1373-2  
**Tobramycin and Dexamethasone Ophthalmic Suspension, USP**  
**0.3%/0.1%**  
 For Topical Ophthalmic Use Only  
 Sterile  
 2.5 mL Rx only

**amneal**

Mfg. Lic. No. G/28/1539  
 Manufactured by:  
**Anneal Pharmaceuticals Pvt. Ltd.**  
 Parenteral Unit  
 Ahmedabad 382213, INDIA  
 Distributed by:  
**Anneal Pharmaceuticals LLC**  
 Bridgewater, NJ 08807  
 Rev. 11-2020-02

  
 N 3 69238 13732 3

NDC 69238-1373-2  
**Tobramycin and Dexamethasone Ophthalmic Suspension, USP**  
**0.3%/0.1%**  
 2.5 mL Rx only



**NDC 69238-1374-3  
 Tobramycin and Dexamethasone Ophthalmic Solution USP, 0.3%/0.1%**

**Rx Only**  
**5 mL**  
**Amneal Pharmaceuticals LLC**

Unvarnished Area for,  
(6 x 14 mm)

NDC 69238-1374-3  
**Tobramycin and  
Dexamethasone  
Ophthalmic  
Suspension, USP**

**0.3%/0.1%**

For Topical Ophthalmic Use Only  
Sterile  
5 mL

Rx only

**Each mL contains:**

**Active:** Tobramycin, USP 0.3% (3 mg) and Dexamethasone, USP 0.1% (1 mg).

**Preservative:** Benzalkonium chloride 0.01%. **Inactive:** Tyloxapol, edetate disodium, sodium chloride, hydroxyethyl cellulose, sodium sulfate, sulfuric acid and/or sodium hydroxide (to adjust pH) and water for injection, USP.

**PRECAUTION:** Do not touch dropper tip to any surface, as this may contaminate the suspension.

**Store at 8° to 27°C (46° to 80°F). Store upright.**

**SHAKE WELL BEFORE USING.**

**Usual Dosage:**

READ ENCLOSED INSERT.

Mfg. Lic. No. G728/1539

Distributed by:

**Amneal Pharmaceuticals LLC**

Bridgewater, NJ 08807

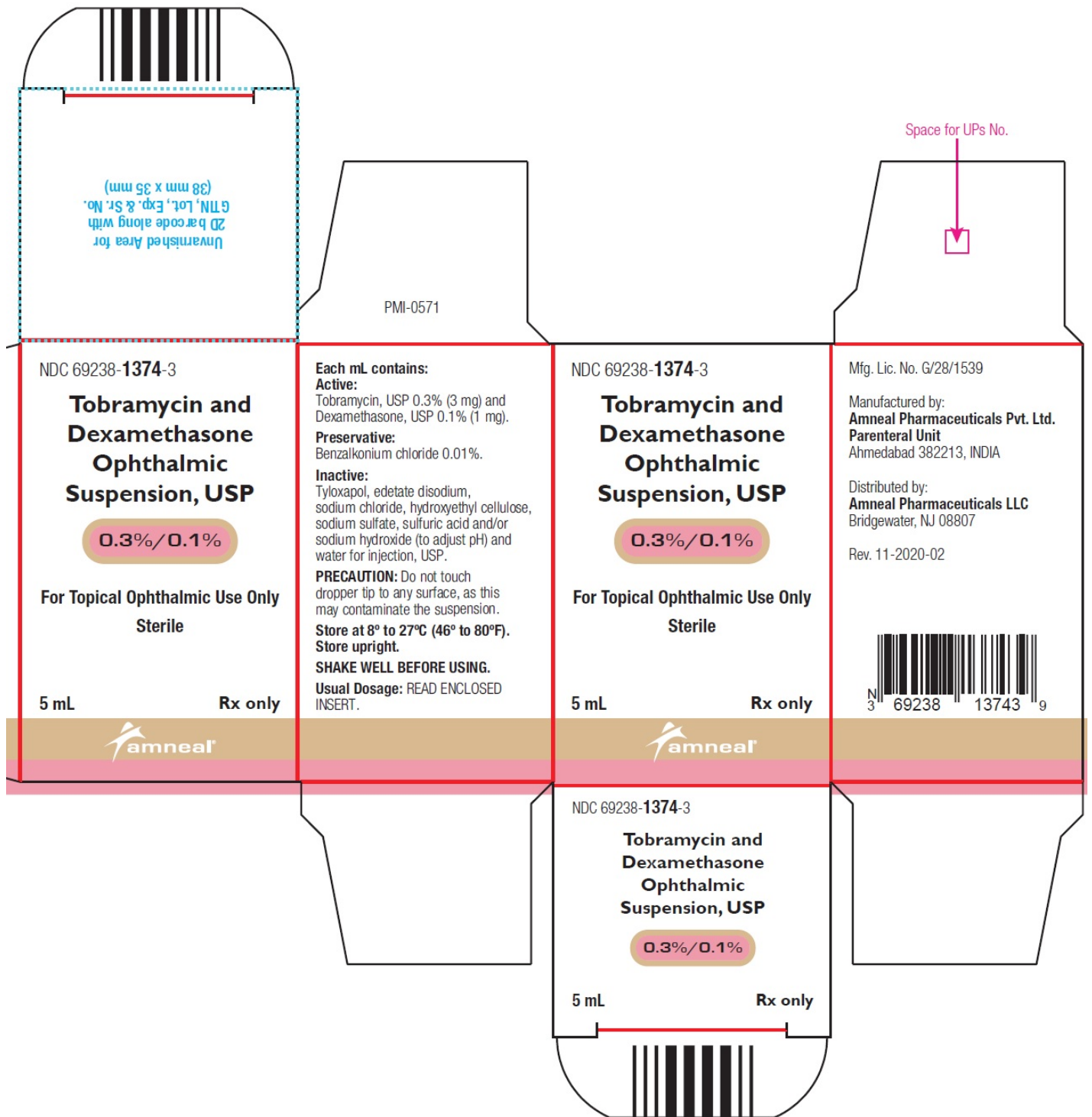
Rev. 04-2024-04 Made in INDIA



(01)\_0\_0369238\_13743\_9

Unvarnished Area for,  
Lot. & Exp.  
(10 x 14 mm)





**NDC 69238-1375-6**  
**Tobramycin and Dexamethasone Ophthalmic Solution USP, 0.3%/0.1%**  
**Rx Only**  
**10 mL**  
**Amneal Pharmaceuticals LLC**

Unvarnished Area for  
(6 mm x 18 mm)

NDC 69238-1375-6  
**Tobramycin and  
Dexamethasone  
Ophthalmic  
Suspension, USP**

0.3%/0.1%

For Topical Ophthalmic Use Only  
Sterile

10 mL

Rx only

**Each mL contains:**  
**Active:** Tobramycin, USP  
0.3% (3 mg) and  
Dexamethasone, USP  
0.1% (1 mg).  
**Preservative:** Benzalkonium  
chloride 0.01%. **Inactive:**  
Tyloxapol, edetate disodium,  
sodium chloride,  
hydroxyethyl cellulose,  
sodium sulfate, sulfuric acid  
and/or sodium hydroxide (to  
adjust pH) and water for  
injection, USP.

**PRECAUTION:** Do not touch  
dropper tip to any surface,  
as this may contaminate  
the suspension.

**Store at 8° to 27°C (46° to  
80°F). Store upright.  
SHAKE WELL BEFORE  
USING.**

**Usual Dosage:**  
READ ENCLOSED INSERT.  
Mfg. Lic. No. G/28/1539  
Distributed by:  
**Anneal Pharmaceuticals LLC**  
Bridgewater, NJ 08807  
Rev. 04-2024-04 Made in INDIA



Unvarnished Area for  
Lot. and Exp.  
(10 mm x 18 mm)

Unvarnished Area for  
2D barcode along with  
GTIN, Lot, Exp. & Sr. No.  
(38 mm x 35 mm)

PMI-0573

Space for UPs No.

NDC 69238-1375-6

**Tobramycin and  
Dexamethasone  
Ophthalmic  
Suspension, USP**

0.3%/0.1%

For Topical Ophthalmic Use Only  
Sterile

10 mL

Rx only



**Each mL contains:**

**Active:**  
Tobramycin, USP 0.3% (3 mg) and  
Dexamethasone, USP 0.1% (1 mg).

**Preservative:**  
Benzalkonium chloride 0.01%.

**Inactive:**  
Tyloxapol, edetate disodium,  
sodium chloride, hydroxyethyl cellulose,  
sodium sulfate, sulfuric acid and/or  
sodium hydroxide (to adjust pH) and  
water for injection, USP.

**PRECAUTION:**  
Do not touch dropper tip to any surface,  
as this may contaminate the suspension.

**Store at 8° to 27°C (46° to 80°F).  
Store upright.**

**SHAKE WELL BEFORE USING.**

**Usual Dosage:** READ ENCLOSED  
INSERT.

NDC 69238-1375-6

**Tobramycin and  
Dexamethasone  
Ophthalmic  
Suspension, USP**

0.3%/0.1%

For Topical Ophthalmic Use Only  
Sterile

10 mL

Rx only



Mfg. Lic. No. G/28/1539

Manufactured by:  
**Anneal Pharmaceuticals Pvt. Ltd.**  
Parenteral Unit  
Ahmedabad 382213, INDIA

Distributed by:  
**Anneal Pharmaceuticals LLC**  
Bridgewater, NJ 08807

Rev. 11-2020-02



NDC 69238-1375-6

**Tobramycin and  
Dexamethasone  
Ophthalmic  
Suspension, USP**

0.3%/0.1%

10 mL

Rx only



# TOBRAMYCIN AND DEXAMETHASONE

tobramycin and dexamethasone suspension/ drops

## Product Information

<b>Product Type</b>	HUMAN PRESCRIPTION DRUG	<b>Item Code (Source)</b>	NDC:69238-1373
<b>Route of Administration</b>	OPHTHALMIC		

## Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>TOBRAMYCIN</b> (UNII: VZ8RRZ51VK) (TOBRAMYCIN - UNII:VZ8RRZ51VK)	TOBRAMYCIN	3 mg in 1 mL
<b>DEXAMETHASONE</b> (UNII: 7S5I7G3JQL) (DEXAMETHASONE - UNII:7S5I7G3JQL)	DEXAMETHASONE	1 mg in 1 mL

## Inactive Ingredients

Ingredient Name	Strength
<b>BENZALKONIUM CHLORIDE</b> (UNII: F5UM2KM3W7)	
<b>EDETATE DISODIUM</b> (UNII: 7FLD91C86K)	
<b>HYDROXYETHYL CELLULOSE (2000 CPS AT 1%)</b> (UNII: S38J6RZN16)	
<b>SODIUM CHLORIDE</b> (UNII: 451W47IQ8X)	
<b>SODIUM HYDROXIDE</b> (UNII: 55X04QC32I)	
<b>SODIUM SULFATE</b> (UNII: 0YPR65R21J)	
<b>SULFURIC ACID</b> (UNII: O40UQP6WCF)	
<b>TYLOXAPOL</b> (UNII: Y27PUL9H56)	
<b>WATER</b> (UNII: 059QF0KO0R)	

## Product Characteristics

<b>Color</b>	white (white to off-white uniform suspension)	<b>Score</b>	
<b>Shape</b>		<b>Size</b>	
<b>Flavor</b>		<b>Imprint Code</b>	
<b>Contains</b>			

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69238-1373-2	1 in 1 CARTON	07/17/2021	
1		2.5 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA212991	07/17/2021	

## TOBRAMYCIN AND DEXAMETHASONE

tobramycin and dexamethasone suspension/ drops

### Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:69238-1374
Route of Administration	OPHTHALMIC		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
TOBRAMYCIN (UNII: VZ8RRZ51VK) (TOBRAMYCIN - UNII:VZ8RRZ51VK)	TOBRAMYCIN	3 mg in 1 mL
DEXAMETHASONE (UNII: 7S5I7G3JQL) (DEXAMETHASONE - UNII:7S5I7G3JQL)	DEXAMETHASONE	1 mg in 1 mL

### Inactive Ingredients

Ingredient Name	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
HYDROXYETHYL CELLULOSE (2000 CPS AT 1%) (UNII: S38J6RZN16)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
SODIUM SULFATE (UNII: 0YPR65R21J)	
SULFURIC ACID (UNII: O40UQP6WCF)	
TYLOXAPOL (UNII: Y27PUL9H56)	
WATER (UNII: 059QF0KO0R)	

### Product Characteristics

Color	white (white to off-white uniform suspension)	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69238-1374-3	1 in 1 CARTON	07/17/2021	
1		5 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA212991	07/17/2021	

## TOBRAMYCIN AND DEXAMETHASONE

tobramycin and dexamethasone suspension/ drops

### Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:69238-1375
Route of Administration	OPHTHALMIC		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>TOBRAMYCIN</b> (UNII: VZ8RRZ51VK) (TOBRAMYCIN - UNII:VZ8RRZ51VK)	TOBRAMYCIN	3 mg in 1 mL
<b>DEXAMETHASONE</b> (UNII: 7S5I7G3JQL) (DEXAMETHASONE - UNII:7S5I7G3JQL)	DEXAMETHASONE	1 mg in 1 mL

### Inactive Ingredients

Ingredient Name	Strength
<b>BENZALKONIUM CHLORIDE</b> (UNII: F5UM2KM3W7)	
<b>EDETATE DISODIUM</b> (UNII: 7FLD91C86K)	
<b>HYDROXYETHYL CELLULOSE (2000 CPS AT 1%)</b> (UNII: S38J6RZN16)	
<b>SODIUM CHLORIDE</b> (UNII: 451W47IQ8X)	
<b>SODIUM HYDROXIDE</b> (UNII: 55X04QC32I)	
<b>SODIUM SULFATE</b> (UNII: 0YPR65R21J)	
<b>SULFURIC ACID</b> (UNII: O40UQP6WCF)	
<b>TYLOXAPOL</b> (UNII: Y27PUL9H56)	
<b>WATER</b> (UNII: 059QF0KO0R)	

### Product Characteristics

Color	white (white to off-white uniform suspension)	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69238-1375-6	1 in 1 CARTON	07/17/2021	
1		10 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA212991	07/17/2021	

**Labeler** - Amneal Pharmaceuticals NY LLC (123797875)

Revised: 8/2024

Amneal Pharmaceuticals NY LLC