

GOODSENSE EYE DROPS ADVANCED RELIEF MOISTURIZER- dextran, polyethylene glycol 400, povidone, tetrahydrozoline hcl solution Preferred PHarmaceuticals Inc.

GoodSense Eye Drops Advanced Relief Moisturizer (PLD)

Active Ingredients

Dextran 70.....0.1%

Polyethylene glycol 400.....1%

Povidone.....1%

tetrahydrozoline HCl.....0.05%

Purposes

Dextran 70.....Lubricant

Polyethylene glycol 400.....Lubricant

Povidone.....Lubricant

Tetrahydrozoline HClRedness reliever

Uses

- relieves redness of the eye due to minor eye irritations
- as a lubricant to prevent further irritation or to relieve dryness of the eye

Warnings

For external use only

Ask a doctor before use if you have narrow angle glaucoma

Stop use and ask a doctor if you experience

- eye pain
- changes in vision
- continued redness or irritation of the eye, or if the condition worsens or persists for more than 72 hours

When using this product

- pupils may become enlarged temporarily
- to avoid contamination, do not touch tip of container to any surface. Replace cap after using.
- if solution changed color or becomes cloudy, do not use
- overuse may produce increased redness of the eye
- remove contact lens before using

If pregnant or breast-feeding , ask a health professional before use.

Keep out of reach of children. If swallowed, get medical help or contact a Poison control Center (1-800-111-1222) right away.

Directions

Instill 1 to 2 drops in the affected eye(s) up to 4 times daily.

Relabeled By: Preferred Pharmaceuticals Inc.

NDC 68788-8635-1

Inactive ingredients

benzalkonium chloride, boric acid, edetate disodium, purified water, sodium borate, and sodium chloride

<p align="center">Eye Drops Advanced Relief Moisturizer</p> <p>Visine® Advanced Relief</p> <p>Active Ingredients: Tetrahydrozoline HCl 0.05 %..... Redness reliever / Dextran 70 0.1%, Polyethylene glycol 400 1%, Povidone1%... Lubricants</p> <p>Pkg Size: Exp Date: ####/#### Lot#: Batch#:</p> <p>Ins: Mfg: K.C. Pharmaceuticals, Inc. Prod#:</p> <p>Warning Keep out of the reach of children. For external use only. Ask a doctor before use if you have narrow angle glaucoma. When using this product pupils may become enlarged temporarily, do not touch tip of container to any surface. Replace cap after using. Remove contact lenses before use. Do not use this product if solution changes color or becomes cloudy. Store at 15° to 30°C (59° to 86°F). Keep tightly closed. See box for uses and drug facts.</p>		<p>CAUTION: Federal law PROHIBITS transfer of this drug to any person other than the patient for whom it was prescribed.</p>	<p>Eye Drops Advanced Relief Moisturizer Qty: Ins: Lot: Bat: Prod# (NDC):</p>	Log
<p>Instill _____ drops every _____ hours.</p>	 <p>Directions English</p>	 <p>GTIN ##### SN ##### EXP #####</p>	<p>Eye Drops Advanced Relief Moisturizer Qty: Ins: Lot: Bat: Prod# (NDC):</p>	Chart
		<p>Instrucciones Espanol: Pongase _____ gota(s) cada _____ horas.</p>	<p>Eye Drops Advanced Relief Moisturizer Qty: Ins: Insurance NDC: Lot: Bat:</p>	Billing
			<p>Eye Drops Advanced Relief Moisturizer Qty: Ins: Lot: Bat: Prod# (NDC):</p>	Patient

GOODSENSE EYE DROPS ADVANCED RELIEF MOISTURIZER

dextran, polyethylene glycol 400, povidone, tetrahydrozoline hcl solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:68788-8635(NDC:50804-130)
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ) (POLYETHYLENE GLYCOL 400 - UNII:B697894SGQ)	POLYETHYLENE GLYCOL 400	1 g in 100 mL
TETRAHYDROZOLINE HYDROCHLORIDE (UNII: 0YZT43HS7D) (TETRAHYDROZOLINE - UNII:S9U025Y077)	TETRAHYDROZOLINE HYDROCHLORIDE	0.05 g in 100 mL
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E) (POVIDONE - UNII:FZ989GH94E)	POVIDONE, UNSPECIFIED	1 g in 100 mL
DEXTRAN 70 (UNII: 7SA290YK68) (DEXTRAN 70 - UNII:7SA290YK68)	DEXTRAN 70	0.1 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	

BORIC ACID (UNII: R5ZHV85D4)	
SODIUM BORATE (UNII: 91MBZ8H3QO)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
WATER (UNII: 059QF0KO0R)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68788-8635-1	1 in 1 CARTON	04/16/2024	
1		15 mL in 1 CARTON; Type 0: Not a Combination Product		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M018	04/16/2024	

Labeler - Preferred PHarmaceuticals Inc. (791119022)

Registrant - Preferred PHarmaceuticals Inc. (791119022)

Establishment			
Name	Address	ID/FEI	Business Operations
Preferred PHarmaceuticals Inc.		791119022	RELABEL(68788-8635)

Revised: 5/2025

Preferred PHarmaceuticals Inc.