

**PHARMASAVE ADVANCED RELIEF EYE DROPS 15ML- dextran 70, povidone, polyethylene glycol 400, tetrahydrozoline hydrochloride solution/ drops
KC Pharmaceuticals, Inc.**

Pharmasave Advanced Relief Eye Drops 15mL (MFR- EXPORT)

Active ingredients

Dextran 70 0.1%

Povidone 1%

Polyethylene glycol 400 1%

Tetrahydrozoline hydrochloride 0.05%

Purposes

Lubricant eye drop

Redness reliever

Warnings

For use in the eye only

Do not use

- if you use a monoamine oxidase inhibitor (MAOI)
- if solution changes colour or becomes cloudy.

Ask a doctor or pharmacist before use if you

- have narrow angle glaucoma
- are pregnant or breastfeeding.

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
- overuse of this product may cause increased redness of the eye.

Stop use and ask a doctor if

- you have a headache, eye pain, changes in vision, redness or irritation of the eye
- symptoms worsen or last longer than 3 days.

Keep out of the reach of children.

If swallowed, call a poison control centre or get medical help right away.

Directions

Adults and children 6 years and over: Remove contact lenses before using. Squeeze 1 or 2 drops in the affected eye(s) up to 4 times a day or as directed by a doctor.

Store between 15°C and 30°C. Do not use if safety seal around cap is missing or broken.

Inactive ingredients

Benzalkonium Chloride, Boric acid, Disodium EDTA, Sodium Borate, Sodium Chloride, Water

Questions?

Call 1-800-268-4127 ext. 3

Pharmasave Advanced Relief Eye Drops 15mL



PHARMASAVE ADVANCED RELIEF EYE DROPS 15ML

dextran 70, povidone, polyethylene glycol 400, tetrahydrozoline hydrochloride solution/ drops

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:55651-990
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
DEXTRAN 70 (UNII: 7SA290YK68) (DEXTRAN 70 - UNII:7SA290YK68)	DEXTRAN 70	0.1 g in 100 mL

POVIDONE (UNII: FZ989GH94E) (POVIDONE, UNSPECIFIED - UNII:FZ989GH94E)	POVIDONE	1 g in 100 mL
TETRAHYDROZOLINE HYDROCHLORIDE (UNII: OYZT43HS7D) (TETRAHYDROZOLINE - UNII:S9U025Y077)	TETRAHYDROZOLINE HYDROCHLORIDE	0.05 g in 100 mL
POLYETHYLENE GLYCOL 400 (UNII: B6978945GQ) (POLYETHYLENE GLYCOL, UNSPECIFIED - UNII:3WJQ0SDW1A)	POLYETHYLENE GLYCOL 400	1 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
BORIC ACID (UNII: R57ZHV85D4)	
EDETATE DISODIUM ANHYDROUS (UNII: 8NLQ36F6MM)	
SODIUM BORATE (UNII: 91MBZ8H3QO)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:55651-990-15	1 in 1 BOX	05/14/2022	
1		15 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
Export only		05/14/2022	

Labeler - KC Pharmaceuticals, Inc. (174450460)

Establishment

Name	Address	ID/FEI	Business Operations
KC Pharmaceuticals, Inc.		174450460	manufacture(55651-990) , pack(55651-990) , label(55651-990)

Revised: 3/2023

KC Pharmaceuticals, Inc.