CVS HYDRATION EYE DROPS- polyetylene glycol 400, propylene glycol solution/ drops
CVS

CVS Hydration Eye Drops 10 mL (PLD)

Active ingredients

Polyethylene glycol 400 0.4%

Propylene glycol 0.3%

Purposes

Lubricant

Lubricant

Use

- for use as a protectant against further irritation or to relieve dryness of the eye

Warnings

For external use only

Do not use

- if this product changes color or becomes cloudy
- if you are sensitive to any ingredient in this product

When using this product

- do not touch the tip of container to any surface to avoid contamination
- replace cap after each use

Stop use and ask a doctor

if you experience any of the following:

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

Keep out of reach of childeren.

If swallowed, get medical help or contact a Poison Control Center (1-800-222-1222) right away.

Directions

put 1 or 2 drops in the affected eye(s) as needed

Other information

Store at temperature, not exceeding 59°F (15°C)

Inactive ingredients

boric acid, hypromellose, potassium chloride, purified water, sodium chloride, sodium hyaluronate, sodium hydroxide

CVS Hydration Eye Drops

CVS HYDRATION EYE DROPS

polyetylene glycol 400, propylene glycol solution/ drops

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:51316-615

Route of Administration	OPHTHALMIC

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
PROPYLENE GLYCOL (UNII: 6DC9Q167V3) (PROPYLENE GLYCOL - UNII:6DC9Q167V3)	PROPYLENE GLYCOL	0.3 g in 100 mL
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ) (POLYETHYLENE GLYCOL 400 - UNII:B697894SGQ)	POLYETHYLENE GLYCOL 400	0.4 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
BORIC ACID (UNII: R57ZHV85D4)	
POTASSIUM CHLORIDE (UNII: 660YQ98I10)	
WATER (UNII: 059QF0KO0R)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
HYALURONATE SODIUM (UNII: YSE9PPT4TH)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:51316-615-10	1 in 1 BOX	01/19/2023	
1		10 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
OTC Monograph Drug	M018	01/19/2023	

Labeler - CVS (062312574)

Registrant - KC Pharmaceuticals, Inc. (174450460)

Establishment

Name	Address	ID/FEI	Business Operations
Unimed		688016567	manufacture(51316-615) , pack(51316-615) , label(51316-615)

Revised: 12/2023 CVS