

CVS MULTI-SYMPTOM EYE DROPS- polyethylene glycol 400, tetrahydrozoline hcl, zinc sulfate solution/ drops
CVS Pharmacy, Inc.

CVS Multi-Symptom Eye Drops 15mL (PLD)

Active ingredients

Polyethylene glycol 400 1%

Tetrahydrozoline HCl 0.05%

Zinc sulfate 0.25%

Purposes

Lubricant

Redness reliever

Astringent

Uses

- relieves dryness of the eyes
- for the temporary relief of discomfort and redness of the eye due to minor eye irritations
- for the temporary relief of burning and irritation due to exposure to wind or sun
- for protection against further irritation

Warnings

For external use only

Ask a doctor before use

if you have narrow angle glaucoma

When using this product

- pupils may become enlarged temporarily
- overuse may cause more eye redness
- remove contact lenses before using
- do not touch tip of container to any surface to avoid contamination
- replace cap after using
- do not use if this solution changes color or becomes cloudy

Stop use and ask a doctor if

- you feel eye pain
- changes in vision occur

- redness or irritation of the eye lasts
- condition worsens or lasts more than 72 hours

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 (1-800-222-1222) right away.

Directions

- instill 1 to 2 drops in the affected eye(s) up to 4 times daily
- children under 6 years of age: ask a doctor

Other information

- some users may experience a brief tingling sensation
- store at 20°-25°C (68°-77°F)

Inactive ingredients

benzalkonium chloride, boric acid, edetate disodium, glycerin, hypromellose, purified water, sodium chloride, sodium citrate

CVS Multi-Symptom Eye Drops 15mL



CVS MULTI-SYMPATOM EYE DROPS

polyethylene glycol 400, tetrahydrozoline hcl, zinc sulfate solution/ drops

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69842-884
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

ZINC SULFATE (UNII: 89DS0H96TB) (ZINC CATION - UNII:13S1S8SF37)			ZINC SULFATE	0.25 g in 100 mL
Inactive Ingredients				
Ingredient Name			Strength	
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)				
BORIC ACID (UNII: R57ZHV85D4)				
GLYCERIN (UNII: PDC6A3C0OX)				
WATER (UNII: 059QF0KO0R)				
SODIUM CITRATE (UNII: 1Q73Q2JULR)				
EDETATE DISODIUM (UNII: 7FLD91C86K)				
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)				
SODIUM CHLORIDE (UNII: 451W47IQ8X)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69842-884-15	1 in 1 BOX	05/16/2022	
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
OTC Monograph Drug		M018	05/16/2022	

Labeler - CVS Pharmacy, Inc. (062312574)

Revised: 12/2023

CVS Pharmacy, Inc.