

CVS GENTLE TEARS- dextran 70, hypromellose solution/ drops
CVS Pharmacy, Inc.

CVS Gentle Tears (PLD)

Active ingredients

Dextran 70 0.1%

Hypromellose 0.3%

Purposes

Lubricant

Lubricant

Use

- temporary relief of discomfort due to minor irritations of the eye or to exposure to wind or sun

Warnings

For external use only

Do not use

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

When using this product

- to avoid contamination, do not touch tip of container to any surface
- do not reuse
- once opened, discard

Stop use and ask a doctor if

- you feel eye pain
- changes in vision occur
- redness or irritation of the eye(s) gets worse 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

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

Other information

store at 15°-25°C (59°-77°F)

Inactive ingredients

boric acid, disodium edetate hydrate, potassium chloride, purified water, sodium borate, sodium chloride

CVS Gentle Tears 35ct



CVS GENTLE TEARS

dextran 70, hypromellose solution/ drops

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69842-474
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO) (HYPROMELLOSE, UNSPECIFIED - UNII:3NXW29V3WO)	HYPROMELLOSE, UNSPECIFIED	0.3 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
EDETATE DISODIUM (UNII: 7FLD91C86K)	
SODIUM BORATE (UNII: 91MBZ8H3QO)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
BORIC ACID (UNII: R57ZHV85D4)	
POTASSIUM CHLORIDE (UNII: 660YQ98I10)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69842-474-01	35 in 1 BOX	01/01/2020	
1		0.4 mL in 1 VIAL, SINGLE-USE; 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	01/01/2020	

Labeler - CVS Pharmacy, Inc. (062312574)

Registrant - KC Pharmaceuticals, Inc. (174450460)

Establishment

Name	Address	ID/FEI	Business Operations
KC Pharmaceuticals, Inc.		174450460	pack(69842-474) , label(69842-474)

Establishment

Name	Address	ID/FEI	Business Operations
Unimed		689852052	manufacture(69842-474)

Revised: 12/2025

CVS Pharmacy, Inc.