

**CVS GENTLE TEARS- dextran 70, hypromellose liquid**  
**CVS Pharmacy, Inc.**

*Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.*

-----

**CVS Gentle Tears**

**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 liquid

### Product Information

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

<b>Active Ingredient/Active Moiety</b>				
<b>Ingredient Name</b>			<b>Basis of Strength</b>	<b>Strength</b>
DEXTRAN 70 (UNII: 7SA290 YK68) (DEXTRAN 70 - UNII:7SA290 YK68)			DEXTRAN 70	0.1 g in 100 mL
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29 V3WO) (HYPROMELLOSE, UNSPECIFIED - UNII:3NXW29 V3WO)			HYPROMELLOSE, UNSPECIFIED	0.3 g in 100 mL
<b>Inactive Ingredients</b>				
<b>Ingredient Name</b>				<b>Strength</b>
BORIC ACID (UNII: R57ZHV85D4)				
POTASSIUM CHLORIDE (UNII: 660YQ98I10)				
WATER (UNII: 059QF0KO0R)				
SODIUM CHLORIDE (UNII: 451W47IQ8X)				
EDETATE DISODIUM (UNII: 7FLD91C86K)				
SODIUM BORATE (UNII: 91MBZ8H3QO)				
<b>Packaging</b>				
#	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		
<b>Marketing Information</b>				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph final	part349	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)