REXALL STERILE ARTIFICIAL TEARS EYE DROPS - glycerin solution
Dolgencorp 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.

Drug Facts

Active ingredients	Purpose
Glycerin 0.2%	Lubricant
Hypromellose 0.2%	Lubricant
Polyethylene glycol 400 1%	Lubricant

Uses
- for the temporary relief of burning and irritation due to dryness of the eye
- for protection against further irritation

Warnings for external use only

When using this product
- remove contact lenses before use
- do not use if this solution changes color or becomes cloudy
- do not touch tip of container or any surface to avoid contamination
- replace cap after each use

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 last 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 right away

Directions
- put 1 to 2 drops in the affected eye(s) as needed.
- children under 6 years of age: ask a doctor.

Other information
- some users may experience a brief tingling sensation
- store at 15° to 25°C (59°-77°F)

Inactive ingredients benzalkonium chloride, boric acid, dextrose, glycine, magnesium chloride, potassium chloride, purified water, sodium borate, sodium chloride, sodium citrate, sodium lactate.

Packaged For:

Dolgencorp, LLC

100 Mission Ridge

Goodlettsville, TN 37072 USA

Made in Korea

REXALL STERILE ARTIFICIAL TEARS EYE DROPS

glycerin solution

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:55910-522
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
GLYCERIN (UNII: PDC6A3C0OX) (GLYCERIN - UNII:PDC6A3C0OX)	GLYCERIN	0.002 mg in 1 mg
HYPROMELLOSES (UNII: 3NXW29V3WO) (HYPROMELLOSES - UNII:3NXW29V3WO)	HYPROMELLOSES	0.002 mg in 1 mg
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ) (POLYETHYLENE GLYCOL 400 - UNII:B697894SGQ)	POLYETHYLENE GLYCOL 400	0.01 mg in 1 mg

Inactive Ingredients	
Ingredient Name	**Strength**
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
BORIC ACID (UNII: R57ZHV85D4)	
DEXTROSE (UNII: IY9XDZ35W2)	
GLYCINE (UNII: TE7660XO1C)	
MAGNESIUM CHLORIDE (UNII: 02F3473H9O)	
POTASSIUM CHLORIDE (UNII: 660YQ98I10)	
WATER (UNII: 059QF0KO0R)	
SODIUM BORATE (UNII: 91MBZ8H3QO)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	
SODIUM LACTATE (UNII: TU7HW0W0QT)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:55910-522-15	1 in 1 CARTON		
1		15 mg in 1 BOTTLE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part349	06/30/2013	

Labeler - Dolgencorp Inc (068331990)

Revised: 6/2013 Dolgencorp Inc