# EXCHANGE SELECT ARTIFICIAL TEARS LUBRICANT EYE DROPS- glycerin, hypromellose, polyethylene glycol 400 solution Army and Air Force Exchange Service

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.

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#### **Exchange Select Artificial Tears**

#### Active ingredients

Glycerin 0.2%

Hypromellose 0.2%

Polyethylene glycol 400 1%

#### **Purpose**

Glycerin....Lubricant

Hypromellose....Lubricant

Polyethylene glycol 400....Lubricant

#### Uses

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

#### Warnings

#### For external use only

**Do not use this product if** solution changes color or becomes cloudy

## Stop use and ask a doctor if you experience

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

#### When using this product

- to avoid contamination, do not touch tip of container to any surface
- replace cap after using. Keep container tightly closed.
- remove contact lens byefore using

**Keep out of reach of children.** If accidentally swallowed, get medical help or contact a Poison Control Center immediately.

### If pregnant or breast feeding, ask a health professional before use

#### Directions

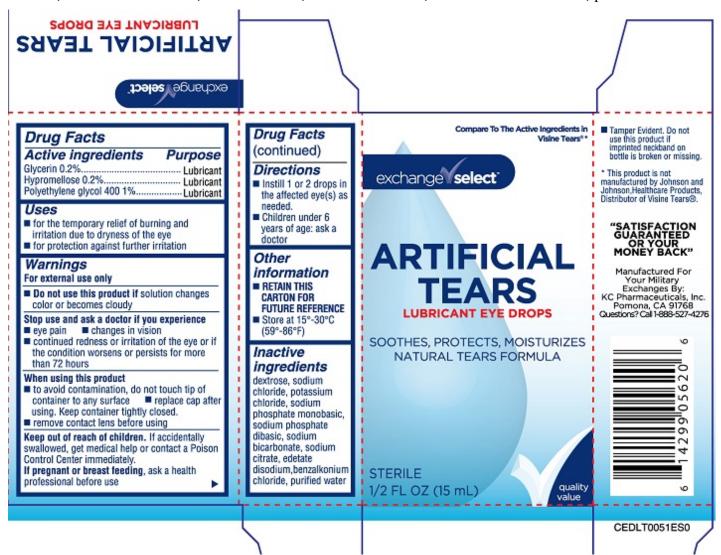
- Instill 1 or 2 drops in the affected eye(s) as needed.
- Children under 6 years of age: ask a doctor

#### Other information

- RETAIN THIS CARTON FOR FUTURE REFERENCE
- Store at 15°-30°C (59°-86°F)

#### **Inactive ingredients**

dextrose, sodium chloride, potassium chloride, sodium phosphate monobasic, sodium phosphate dibasic, sodium bicarbonate, sodium citrate, edetate disodium, benzalkonium chloride, purified water.



#### EXCHANGE SELECT ARTIFICIAL TEARS LUBRICANT EYE DROPS

glycerin, hypromellose, polyethylene glycol 400 solution

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:55301-178
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29 V3WO) (HYPROMELLOSE, UNSPECIFIED - UNII:3NXW29 V3WO)	HYPROMELLOSE, UNSPECIFIED	0.2 g in 100 mL	
GLYCERIN (UNII: PDC6A3C0OX) (GLYCERIN - UNII:PDC6A3C0OX)	GLYCERIN	0.2 g in 100 mL	
<b>POLYETHYLENE GLYCOL 400</b> (UNII: B697894SGQ) (POLYETHYLENE GLYCOL 400 - UNII:B697894SGQ)	POLYETHYLENE GLYCOL 400	1 g in 100 mL	

Inactive Ingredients				
Ingredient Name	Strength			
SODIUM BICARBONATE (UNII: 8 MDF5 V39 QO)				
EDETATE DISO DIUM (UNII: 7FLD9 1C8 6 K)				
SO DIUM PHO SPHATE, DIBASIC (UNII: GR686LBA74)				
DEXTRO SE (UNII: IY9 XDZ35W2)				
SODIUM CHLORIDE (UNII: 451W47IQ8X)				
SODIUM PHO SPHATE, MO NO BASIC (UNII: 3980 JIH2SW)				
SODIUM CITRATE (UNII: 1Q73Q2JULR)				
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)				
WATER (UNII: 059QF0KO0R)				
POTASSIUM CHLORIDE (UNII: 660 YQ98 I10)				

Packaging				
# Item Code	Package Description	Marketing Start Date	Marketing End Date	
1 NDC:55301-178- 01	1 in 1 CARTON	02/12/2020		
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 final	part349	02/12/2020		

# Labeler - Army and Air Force Exchange Service (001695568)

# **Registrant** - K.C. Pharmaceuticals, Inc. (174450460)

Establishment				
Name	Address	ID/FEI	Business Operations	
K.C. Pharmaceuticals, Inc.		174450460	manufacture(55301-178), pack(55301-178), label(55301-178)	

Revised: 2/2020 Army and Air Force Exchange Service