

**ARTIFICIAL TEARS- glycerin solution/ drops  
Preferred Pharmaceuticals 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.*

-----  
**GC Artificial Tears 181**

**Active ingredients**

Glycerin 0.2%

Hypromellose 0.2%

Polyethylene glycol 400 1%

**Purpose**

Lubricant

**Uses**

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

**Warnings**

**For external use only**

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

**When using this product**

- to avoid contamination, do not touch tip of container to any surface. Replace cap after using.
- remove contact lenses before using

**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

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

## Other information

- store at 15°-30°C (59°-86°F)

## Inactive ingredients

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

## Questions or comments?

**1-800-540-3765**

## package label

**Artificial Tears Lubricant Eye Drops**  
Visine®

Active Ingredients Glycerin 0.2%...lubricant  
Hypromellose 0.2%...lubricant Polyethylene glycol 400 1%...lubricant

Pkg Size: Exp Date:  
Lot#: Ins:  
Batch#: Ins:  
Mfg: Geri-Care; Brooklyn, New York  
Prod#: Ins:  
Warning

Keep out of the reach of children. For external use only. When using this product do not touch tip of container to any surface. Replace cap after using. Remove contact lenses before use. Do not use this product if solution changes color or becomes cloudy. If pregnant or breast feeding, ask a health care professional before use. Stop use and ask a doctor if you experience eye pain, changes in vision, conjunctival redness, or irritation of the eye. Store at 15° to 30°C (59° to 86°F). Keep tightly closed. See box for uses and drug facts.

**Directions English**  
Instill \_\_\_ drops every \_\_\_ hours.

**Instrucciones Espanol:**  
Pongase \_\_\_ gota(s) cada \_\_\_ horas.

CAUTION: Federal law PROHIBITS transfer of this drug to any person other than the patient for whom it was prescribed

Artificial Tears Lubricant Eye Drops  
Qty: Ins:  
Lot#: Bat#:  
Prod# (NDC):

Artificial Tears Lubricant Eye Drops  
Qty: Ins:  
Lot#: Bat#:  
Prod# (NDC):

Artificial Tears Lubricant Eye Drops  
Qty: Ins:  
Insurance NDC:  
Lot#: Bat#:

Artificial Tears Lubricant Eye Drops  
Qty: Ins:  
Lot#: Bat#:  
Prod# (NDC):

Log  
Chart  
Billing  
Patient

## ARTIFICIAL TEARS

glycerin solution/ drops

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:68788-7266(NDC:57896-181)
<b>Route of Administration</b>	OPHTHALMIC		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>GLYCERIN</b> (UNII: PDC6A3C0OX) (GLYCERIN - UNII:PDC6A3C0OX)	GLYCERIN	0.002 mg in 1 mg
<b>HYPROMELLOSE, UNSPECIFIED</b> (UNII: 3NXW29V3WO) (HYPROMELLOSES - UNII:3NXW29V3WO)	HYPROMELLOSE, UNSPECIFIED	0.002 mg in 1 mg
<b>POLYETHYLENE GLYCOL 400</b> (UNII: B697894SGQ) (POLYETHYLENE GLYCOL 400 - UNII:B697894SGQ)	POLYETHYLENE GLYCOL 400	0.01 mg in 1 mg

### Inactive Ingredients

Ingredient Name	Strength
<b>DEXTROSE, UNSPECIFIED FORM</b> (UNII: IY9XDZ35W2)	
<b>SODIUM PHOSPHATE, DIBASIC, DIHYDRATE</b> (UNII: 94255I6E2T)	
<b>SODIUM BICARBONATE</b> (UNII: 8MDF5V39QO)	
<b>EDETATE DISODIUM</b> (UNII: 7FLD91C86K)	
<b>BENZALKONIUM CHLORIDE</b> (UNII: F5UM2KM3W7)	
<b>SODIUM PHOSPHATE, MONOBASIC, UNSPECIFIED FORM</b> (UNII: 3980JH2SW)	
<b>POTASSIUM CHLORIDE</b> (UNII: 660YQ98I10)	
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>SODIUM CHLORIDE</b> (UNII: 451W47IQ8X)	
<b>SODIUM CITRATE, UNSPECIFIED FORM</b> (UNII: 1Q73Q2JULR)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68788-7266-0	1 in 1 CARTON	03/11/2021	
1		15 mg in 1 BOTTLE; 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	03/11/2021	

**Labeler** - Preferred Pharmaceuticals Inc. (791119022)

**Registrant** - Preferred Pharmaceuticals Inc. (791119022)

### Establishment

Name	Address	ID/FEI	Business Operations
Preferred Pharmaceuticals Inc.		791119022	RELABEL(68788-7266)