

**HARRIS TEETER EYE DROPS ARTIFICIAL TEARS- glycerin, hypromellose, polyethylene glycol 400 solution/ drops
Harris Teeter, LLC.**

Harris Teeter Eye Drops Artificial Tears 5mL (PLD)

Active ingredients

Glycerin 0.2%

Hypromellose 0.2%

Polyethylene glycol 400 1%

Purpose

Lubricant

Lubricant

Lubricant

Uses

- for protection against further irritation
- for the 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 lens 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

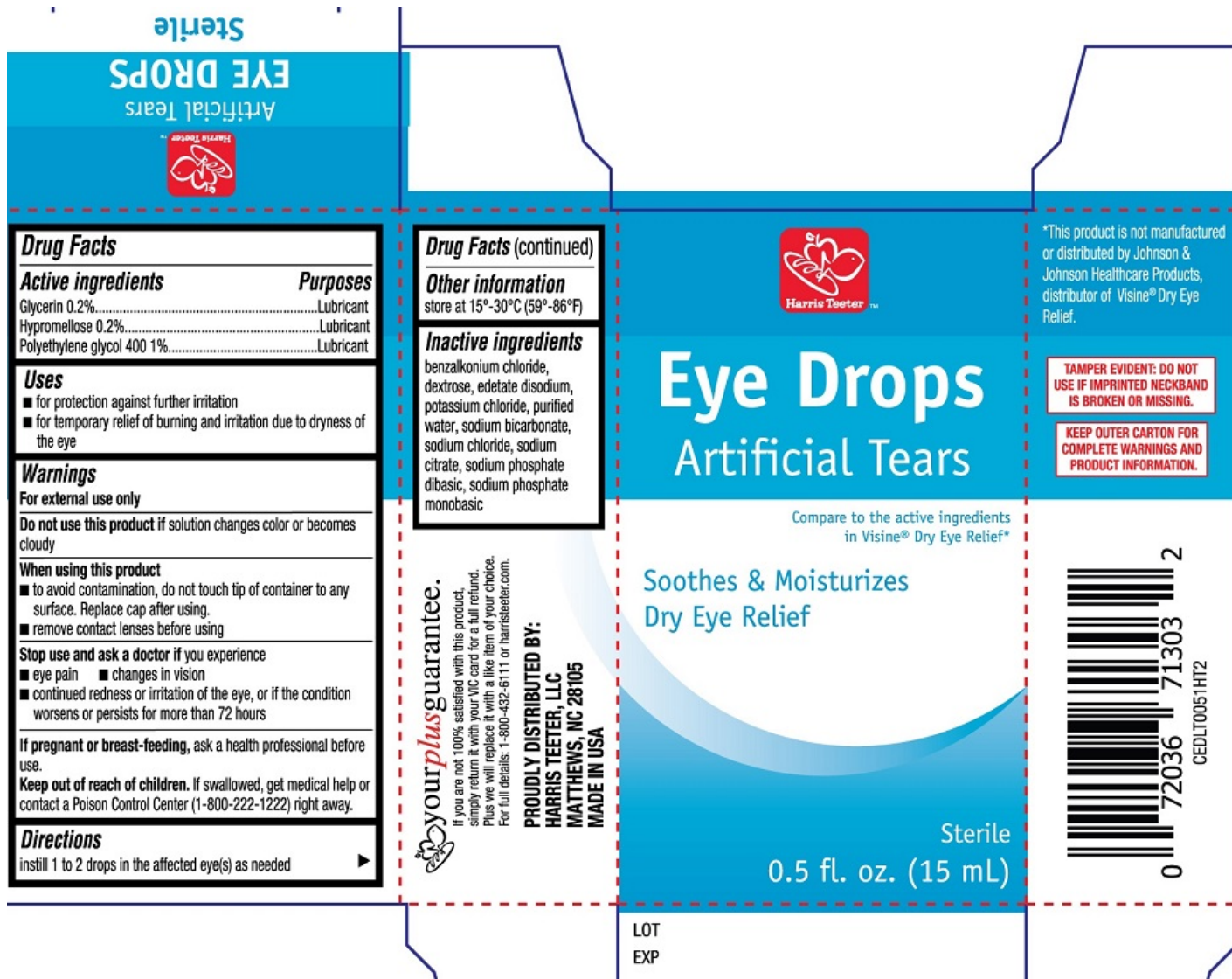
Other information

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

Harris Teeter Eye Drops Artificial Tears



HARRIS TEETER EYE DROPS ARTIFICIAL TEARS

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Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69256-823
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
GLYCERIN (UNII: PDC6A3C0OX) (GLYCERIN - UNII:PDC6A3C0OX)	GLYCERIN	0.2 g in 100 mL
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO) (HYPROMELLOSE, UNSPECIFIED - UNII:3NXW29V3WO)	HYPROMELLOSE, UNSPECIFIED	0.2 g in 100 mL
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ) (POLYETHYLENE GLYCOL 400 - UNII:B697894SGQ)	POLYETHYLENE GLYCOL 400	1 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
DEXTROSE (UNII: IY9XDZ35W2)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
POTASSIUM CHLORIDE (UNII: 660YQ98I10)	
WATER (UNII: 059QF0KO0R)	
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	
SODIUM PHOSPHATE, DIBASIC (UNII: GR686LBA74)	
SODIUM PHOSPHATE, MONOBASIC (UNII: 3980JIH2SW)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69256-823-01	1 in 1 BOX	03/30/2023	
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 Drug	M018	04/30/2019	

Labeler - Harris Teeter, LLC. (048463103)

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

Establishment

Name	Address	ID/FEI	Business Operations
K.C. Pharmaceuticals, Inc.		174450460	manufacture(69256-823) , pack(69256-823) , label(69256-823)

Revised: 12/2023

Harris Teeter, LLC.