EQUATE EYE DROPS DRY EYE RELIEF- glycerin, hypromellose, polyethylene glycol 400 solution/ drops
Wal-Mart Stores, Inc.

Equate Dry Eye Relief Eye Drops 15mL twin pk (PLD)

Active ingredients

Glycerin 0.2%

Hypromellose 0.2%

Polyethylene glycol 400 1%

Purposes

Lubricant

Lubricant

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 to 2 drops in the affected eye(s) as needed

Other information

store at 15°-30°C (59°-86°F)

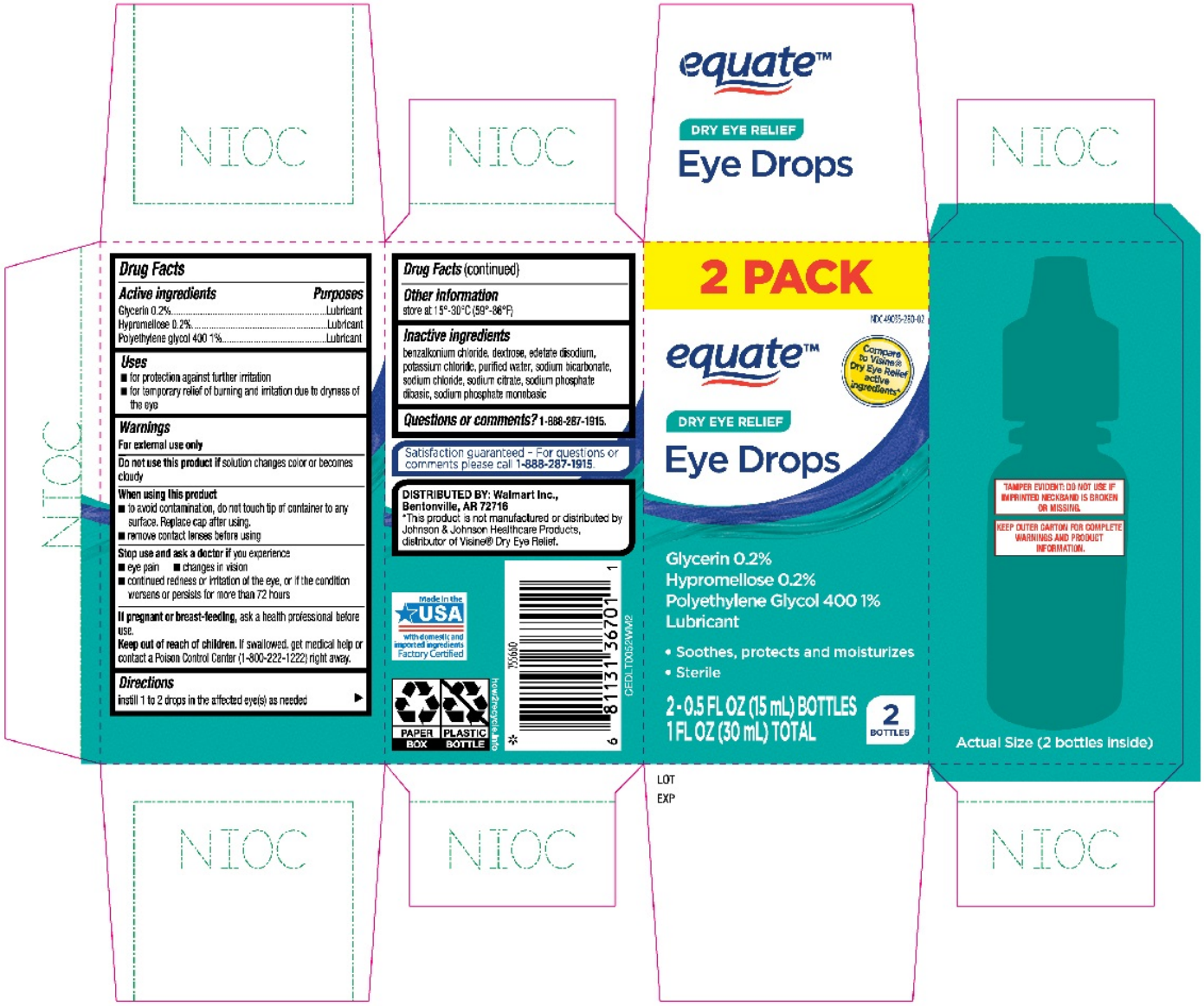
Inactive ingredients

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

Questions or comments?

1-888-287-1915

Equate Dry Eye Relief Eye Drops 2-0.5oz bottles



Equate Eye Drops Dry Eye Relief 2-0.5oz bottles

EQUATE EYE DROPS DRY EYE RELIEF

glycerin, hypromellose, polyethylene glycol 400 solution/ drops

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:49035-280
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, UNSPECIFIED - UNII:3WQ0SDW1A)	POLYETHYLENE GLYCOL 400	1 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
SODIUM PHOSPHATE, MONOBASIC, ANHYDROUS (UNII: KH7I04HPUU)	
POTASSIUM CHLORIDE (UNII: 660YQ98I10)	
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
SODIUM PHOSPHATE DIBASIC DIHYDRATE (UNII: 94255I6E2T)	
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
DEXTROSE (UNII: IY9XDZ35W2)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
WATER (UNII: 059QF0KO0R)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:49035-280-01	2 in 1 BOX	11/01/2019	
1		15 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product		
2	NDC:49035-280-02	2 in 1 BOX	02/01/2021	
2		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	11/01/2019	

Labeler - Wal-Mart Stores, Inc. (051957769)

Registrant - KC Pharmaceuticals, Inc. (174450460)

Establishment

Name	Address	ID/FEI	Business Operations
KC Pharmaceutical, Inc.		174450460	manufacture(49035-280) , pack(49035-280) , label(49035-280)

Revised: 12/2023 Wal-Mart Stores, Inc.