EQUATE ARTIFICIAL TEARS- polyvinyl alcohol, povidone solution/ drops
Walmart, Inc.

Equate Artificial Tears 15mL (PLD)

Active ingredients

Polyvinyl alcohol 0.5%

Povidone 0.6%

Purpose

Lubricant

Lubricant

Uses

- for use as a protectant against further irritation or to relieve dryness of the eye
- for the temporary relief of discomfort due to minor irritations of the eye, or to exposure to wind or sun

Warnings

For external use only

Do not use product if

- solution changes color or becomes cloudy

When using this product

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

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

Keep out of the reach of children.

If accidentally swallowed, get medical help or contact a Poison Control Center (1-800-222-1222) immediately.

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

Questions or comments?

Call 1-888-287-1915

Equate Artificial Tears 15mL

EQUATE ARTIFICIAL TEARS

polyvinyl alcohol, povidone solution/ drops

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:79903-157
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
POVIDONE (UNII: FZ989GH94E) (POVIDONE, UNSPECIFIED - UNII:FZ989GH94E)	POVIDONE	0.6 g in 100 mL
POLYVINYL ALCOHOL, UNSPECIFIED (UNII: 532B59J990) (POLYVINYL	POLYVINYL ALCOHOL,	0.5 g

ALCOHOL, UNSPECIFIED - UNII:532B59J990)	UNSPECIFIED	in 100 mL

Inactive Ingredients

Ingredient Name	Strength
EDETATE DISODIUM (UNII: 7FLD91C86K)	
DEXTROSE (UNII: IY9XDZ35W2)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	
SODIUM PHOSPHATE, DIBASIC, ANHYDROUS (UNII: 22ADO53M6F)	
POTASSIUM CHLORIDE (UNII: 660YQ98I10)	
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
WATER (UNII: 059QF0KO0R)	
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
SODIUM PHOSPHATE, MONOBASIC, ANHYDROUS (UNII: KH7I04HPUU)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:79903-157-05	1 in 1 BOX	02/16/2023	
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	02/16/2023	

Labeler - Walmart, Inc. (051957769)

Registrant - KC Pharmaceuticals, Inc. (174450460)

Establishment

Name	Address	ID/FEI	Business Operations
KC Pharmaceuticals, Inc.		174450460	manufacture(79903-157) , pack(79903-157) , label(79903-157)

Revised: 12/2025 Walmart, Inc.