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

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#### **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

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Product Type HUMAN OTC DRUG Item Code (Source) NDC:79903-157

Route of Administration OPHTHALMIC

Active Ingredient/Active Moiety			
Ingredient Name	<b>Basis of Strength</b>	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	

Inactive Ingredients		
Ingredient Name	Strength	
EDETATE DISODIUM (UNII: 7FLD91C86K)		
DEXTROSE (UNII: IY9XDZ 35W2)		
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)		

l	Packaging			
	# Item Code	Package Description	Marketing Start Date	Marketing End Date
	<b>1</b> 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: 12/2023 Walmart, Inc.