EQUATE RESTORE TEARS- carboxymethylcellulose sodium solution/ drops
Wal-Mart Stores, 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.

Equate Restore Tears Drops

Equate

Restore Tears Drops

49035-189-49

Drug Facts

Active ingredient

Carboxymethylcellulose Sodium 0.5%

Purpose

Eye Lubricant

Uses

● For the temporary relief of burning, irritation, and discomfort due to dryness of the eye or exposure to wind or sun.

● May be used as a protectant against further irritation.

Warnings

For use in the eyes only.

● Retain outer carton for full product drug facts

Do not use

● If solution changes color, or becomes cloudy.

When using this product

● avoid contamination, do not touch tip of container to any surface.

● replace cap after each use.

Stop use and ask a doctor if

● you experience eye pain, changes in vision, continued redness or irritation of the eye(s)

● the condition worsens or persists for more than 72 hours.

Keep out of the reach of children.

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

Directions

● Instill 1or 2 drops into the affected eye(s) as needed.

Other Information

● store at room temperature 15º -30°C (59°-86°F).

● keep tightly closed.

Inactive ingredients

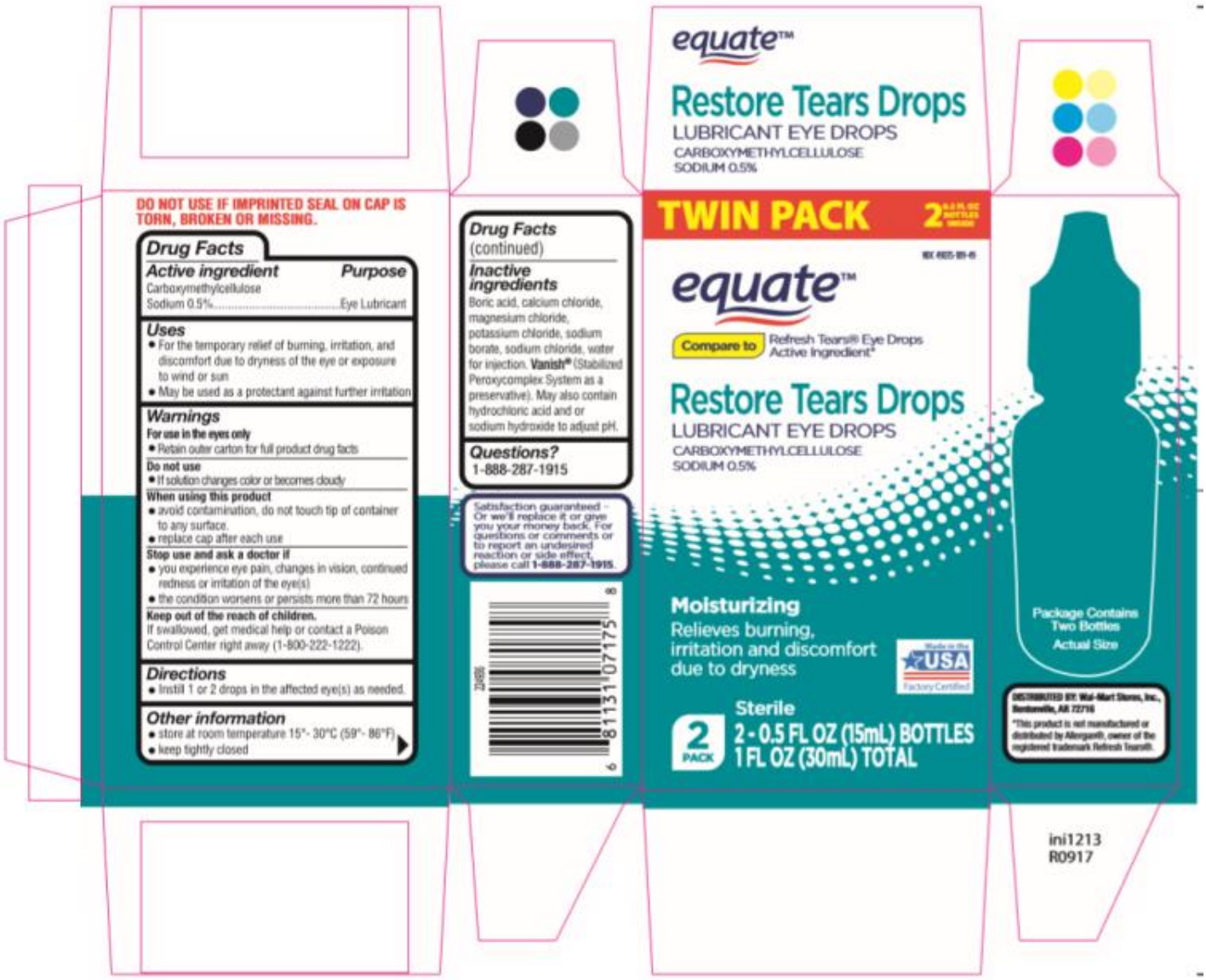
Boric acid, calcium chloride, magnesium chloride, potassium chloride, sodium borate, sodium chloride, water for injection. Vanish® (Stabilized Peroxycomplex System as a preservative). May also contain hydrochloric acid and or sodium hydroxide to adjust pH.

Questions?

1-888-287-1915

PRINCIPAL DISPLAY PANEL

NDC 49035-189-49
equate
Restore Tears Drops
LUBRICANT EYE DROPS
CARBOXYMETHYCELLULOSE SODIUM 0.5%
Sterile
2 – 0.5 fl oz (15mL) BOTTLES
1 FL OZ (30mL) TOTAL



EQUATE RESTORE TEARS

carboxymethylcellulose sodium solution/ drops

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:49035-189
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
CARBOXYMETHYLCELLULOSE SODIUM (UNII: K679OBS311) (CARBOXYMETHYLCELLULOSE - UNII:05JZI7B19X)	CARBOXYMETHYLCELLULOSE SODIUM	5 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
BORIC ACID (UNII: R57ZHV85D4)	

CALCIUM CHLORIDE (UNII: M4I0D6VV5M)	
MAGNESIUM CHLORIDE (UNII: 02F3473H9O)	
POTASSIUM CHLORIDE (UNII: 660YQ98I10)	
SODIUM BORATE (UNII: 91MBZ8H3QO)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
WATER (UNII: 059QF0KO0R)	
CHLORINE DIOXIDE (UNII: 8061YMS4RM)	
POLIHEXANIDE (UNII: 322U039GMF)	

Other Ingredients

Ingredient Kind	Ingredient Name	Quantity
May contain	**HYDROCHLORIC ACID** (UNII: QTT17582CB)	
May contain	**SODIUM HYDROXIDE** (UNII: 55X04QC32I)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:49035-189-49	2 in 1 CARTON	04/10/2014	
1		15 mL 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	04/10/2014	

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

Registrant - Altaire Pharmaceuticals, Inc. (786790378)

Revised: 1/2018 Wal-Mart Stores, Inc.