EQUATE COMFORT- carboxymethylcellulose sodium and hypromelloses gel
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 Comfort Gel

EQUATE COMFORT GEL

49035-197-49

Drug Facts

Active ingredients

Carboxymethylcellulose Sodium 0.25%

Hypromellose 0.3%

Purpose

Eye Lubricant

Eye Lubricant

Uses

● relieves dryness of the eye(s).

● for the temporary relief of discomfort due to minor irritations of the eye from exposure to wind or sun.

● 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 this product 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-30ºC (59º-86ºF).

● keep tightly closed.

Inactive Ingredients:

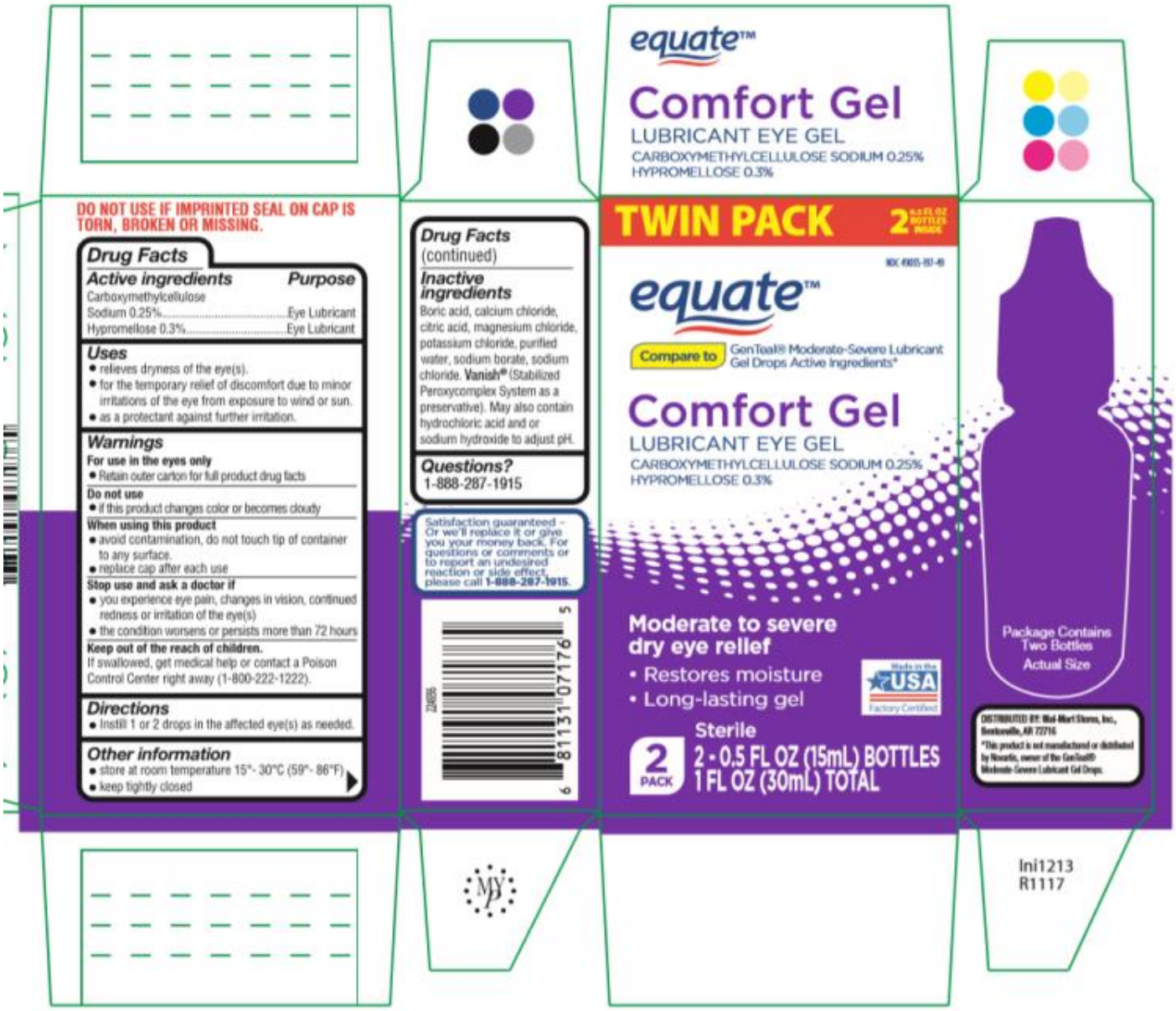
Boric acid, calcium chloride, citric acid, magnesium chloride, potassium chloride, purified water, sodium borate, sodium chloride. Vanish® (Stabilized Peroxycomplex System as a preservative). May also contain hydrochloric acid and or sodium hydroxide to adjust pH.

Questions?

1-888-287-1915

PRINCIPAL DISPLAY PANEL

NDC 49035-197-49
equate
Comfort Gel
LUBRICANT EYE GEL
CARBOXYMETHYCELLULOSE SODIUM 0.25%
HYPROMELLOSE 0.3%
Sterile
2- 0.5 FL OZ (15mL) BOTTLES
1 FL OZ (30mL) TOTAL



EQUATE COMFORT

carboxymethylcellulose sodium and hypromelloses gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:49035-197
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
HYPROMELLOSES (UNII: 3NXW29V3WO) (HYPROMELLOSES - UNII:3NXW29V3WO)	HYPROMELLOSES	3 mg in 1 mL
CARBOXYMETHYLCELLULOSE SODIUM (UNII: K679OBS311) (CARBOXYMETHYLCELLULOSE - UNII:05JZI7B19X)	CARBOXYMETHYLCELLULOSE SODIUM	2.5 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
BORIC ACID (UNII: R57ZHV85D4)	
CALCIUM CHLORIDE (UNII: M4I0D6VV5M)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
MAGNESIUM CHLORIDE (UNII: 02F3473H9O)	
POTASSIUM CHLORIDE (UNII: 660YQ98I10)	
WATER (UNII: 059QF0KO0R)	
SODIUM BORATE (UNII: 91MBZ8H3QO)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
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:49035-197-49	2 in 1 CARTON	04/10/2014	
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 FINAL	part349	04/10/2014	

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

Registrant - Altaire Pharmaceuticals, Inc. (786790378)

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