ROHTO HYDRA- hydroxyethyl cellulose liquid
The Mentholatum Company

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.

Drug Facts

Active ingredient

Hydroxyethyl cellulose 0.6%

Purpose

Hydroxyethyl cellulose - Lubricant

Uses

- temporarily relieves discomfort due to minor irritations of the eye or exposure to wind or sun
- lubricates to prevent further irritation or to relieve dryness of the eye

Warnings

For external use only

When using this product

- do not touch tip of container to any surface to avoid contamination
- replace cap after each use
- do not use if solution changes color or becomes cloudy
- remove contact lenses before using

Stop use and ask a doctor if

- you feel eye pain
- changes in vision occur
- redness or irritation of the eyes lasts
- condition worsens or lasts more than 72 hours

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center right away.

Directions

- put 1 or 2 drops in the affected eye(s) as needed

Other information

- store at 20-25 °C (68-77°F)

- tightly snap on cap to seal

Inactive ingredients

benzalkonium chloride, boric acid, edetate disodium, menthol, polysorbate 80, potassium chloride, purified water, sodium borate, sodium chloride

Questions?

Toll free **1-877-636-2677** MON-FRI 9AM to 5PM (EST)

Principal Display Panel

ROHTO HYDRA			
hydroxyethyl cellulose liquid			
Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:10742-8148
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
HYDROXYETHYL CELLULOSE (2000 MPA.S AT 1%) (UNII: S38J6RZN16) (HYDROXYETHYL CELLULOSE (2000 MPA.S AT 1%) - UNII:S38J6RZN16)	HYDROXYETHYL CELLULOSE (2000 MPA.S AT 1%)	6 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
BORIC ACID (UNII: R57ZHV85D4)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
MENTHOL, UNSPECIFIED FORM (UNII: L7T10EIP3A)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
POTASSIUM CHLORIDE (UNII: 660YQ98I10)	
WATER (UNII: 059QF0KO0R)	
SODIUM BORATE (UNII: 91MBZ8H3QO)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:10742-8148-1	1 in 1 CARTON	01/03/2011	10/31/2018
1		13 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	01/03/2011	10/31/2018

Labeler - The Mentholatum Company (002105757)

Revised: 7/2017 The Mentholatum Company