ROHTO SUN AND SPORT- hypromellose, povidone liquid
The Mentholatum Company

Drug Facts

Active ingredient

Hypromellose 0.3%

Povidone 0.5%

Purpose

Hypromellose – Lubricant

Povidone - Lubricant

Uses

- for the temporary relief of discomfort due to minor irritations of the eye or to exposure to wind or sun
- protects against 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

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 persists for 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
- tightly snap on cap to seal

Other information

- do not store above 25 ^{O}C (77 ^{O}F)

Inactive ingredients

alginic acid, anhydrous citric acid, boric acid, camphor, chlorobutanol, edetate disodium, menthol, polysorbate 80, purified water, sodium borate, sodium citrate, taurine, zinc sulfate

Questions?

1-877-636-2677 MON-FRI 9AM-5PM (EST)

Principal Display Panel

ROHTO SUN AND SPORT
hypromellose, povidone liquid

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:10742-8140
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO) (HYPROMELLOSE, UNSPECIFIED - UNII:3NXW29V3WO)	HYPROMELLOSE, UNSPECIFIED	3 mg in 1 mL
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E) (POVIDONE, UNSPECIFIED - UNII:FZ989GH94E)	POVIDONE, UNSPECIFIED	5 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
ALGINIC ACID (UNII: 8C3Z4148WZ)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
BORIC ACID (UNII: R57ZHV85D4)	
CAMPHOR (SYNTHETIC) (UNII: 5TJD82A1ET)	
CHLOROBUTANOL (UNII: HM4YQM8WRC)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
RACEMENTHOL (UNII: YS08XHA860)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
WATER (UNII: 059QF0KO0R)	
SODIUM BORATE (UNII: 91MBZ8H3QO)	
SODIUM CITRATE, UNSPECIFIED FORM (UNII: 1Q73Q2JULR)	
TAURINE (UNII: 1EQV5MLY3D)	
ZINC SULFATE, UNSPECIFIED FORM (UNII: 89DS0H96TB)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:10742-8140-1	1 in 1 CARTON	07/01/2022	
1		18 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	07/01/2022	

Labeler - The Mentholatum Company (002105757)