ROHTO DUAL LIGHT RELIEF- 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

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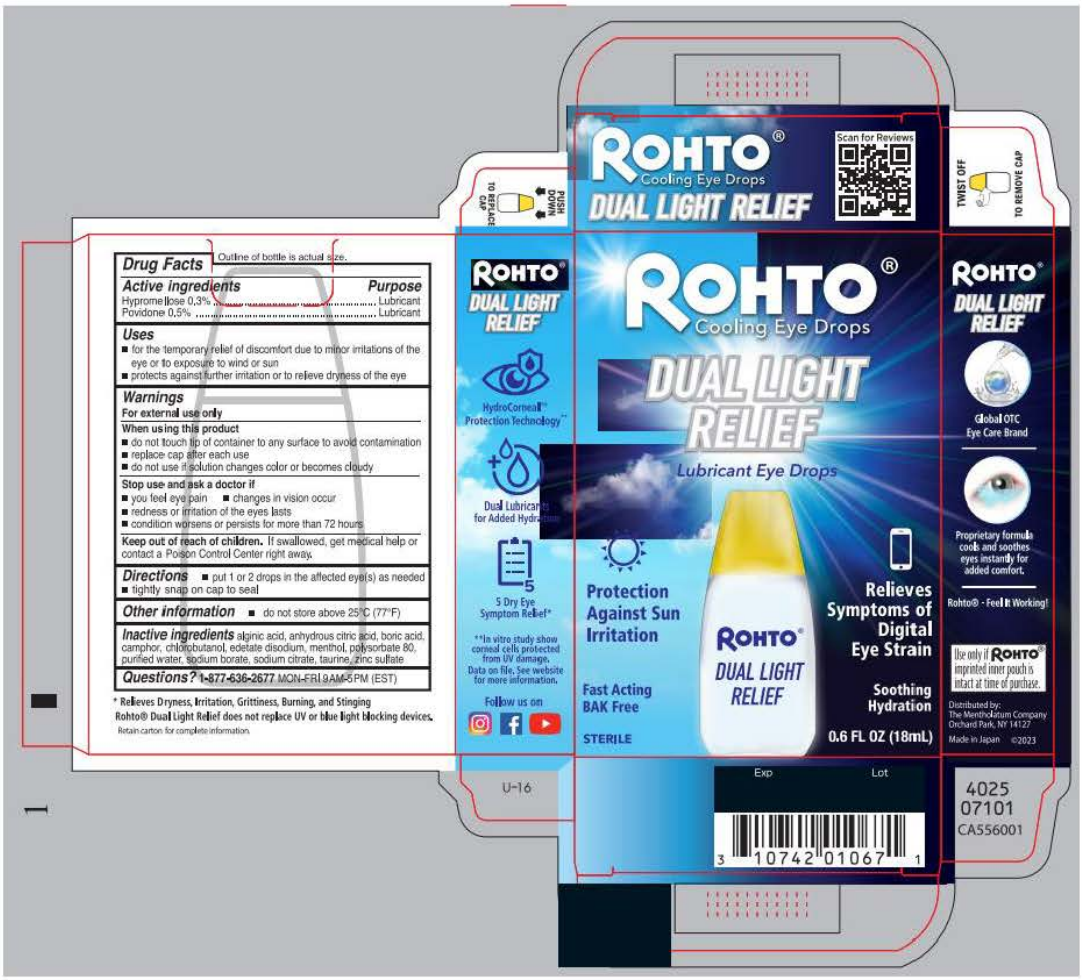

ROHTO DUAL LIGHT RELIEF

hypromellose, povidone liquid

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:10742-8459
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety

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

Inactive Ingredients

Ingredient Name	Strength
LEVOMENTHOL (UNII: BZ1R15MTK7)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
BORIC ACID (UNII: R57ZHV85D4)	
CHLOROBUTANOL (UNII: HM4YQM8WRC)	
TRISODIUM CITRATE DIHYDRATE (UNII: B22547B95K)	
ZINC SULFATE HEPTAHYDRATE (UNII: N57JI2K7WP)	
ALGINIC ACID (UNII: 8C3Z4148WZ)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
SODIUM BORATE (UNII: 91MBZ8H3QO)	
TAURINE (UNII: 1EQV5MLY3D)	
WATER (UNII: 059QF0KO0R)	
CAMPHOR (SYNTHETIC) (UNII: 5TJD82A1ET)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:10742-8459-1	1 in 1 CARTON	05/30/2024	
1		18 mL in 1 BOTTLE, DISPENSING; 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	05/30/2024	

Labeler - The Mentholatum Company (002105757)

Registrant - The Mentholatum Company (002105757)

Establishment

Name	Address	ID/FEI	Business Operations
The Mentholatum Company		002105757	label(10742-8459)

Establishment

Name	Address	ID/FEI	Business Operations
Rohto Pharmaceutical Co., Ltd.		696604024	manufacture(10742-8459)

Revised: 6/2024

The Mentholatum Company