

**ROHTO DUAL LIGHT RELIEF- hypromellose, povidone liquid**  
**Rohto Pharmaceutical Co., Ltd.**

-----  
**Drug Facts - Rohto Dual Light Relief**

**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 OC (77 OF)

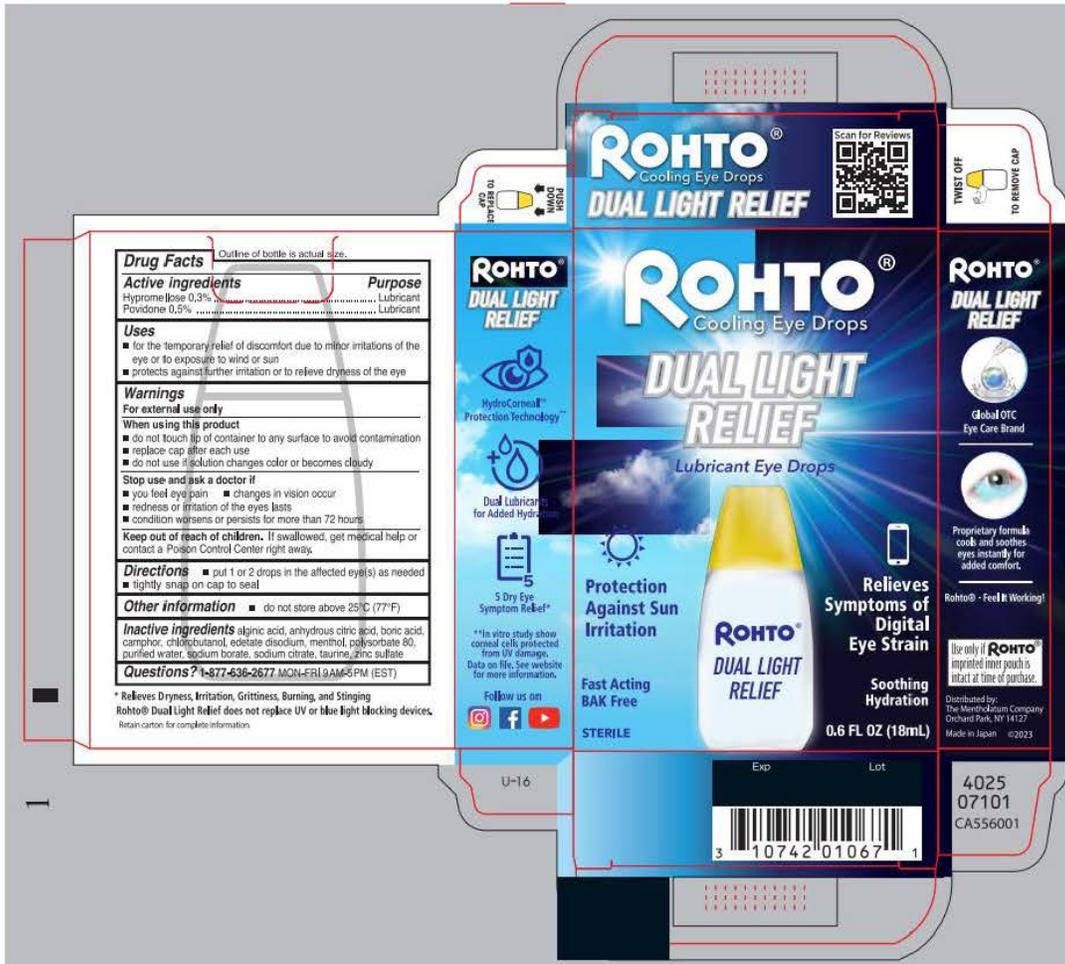
### **Inactive ingredients**

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 DUAL LIGHT RELIEF**

hypromellose, povidone liquid

## Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:66613-8459
<b>Route of Administration</b>	OPHTHALMIC		

## Active Ingredient/Active Moiety

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

## Inactive Ingredients

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

## Packaging

<b>#</b>	<b>Item Code</b>	<b>Package Description</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
1	NDC:66613-8459-1	1 in 1 CARTON	05/30/2024	
1		18 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product		

## Marketing Information

<b>Marketing Category</b>	<b>Application Number or Monograph Citation</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
OTC Monograph Drug	M018	05/30/2024	

**Labeler** - Rohto Pharmaceutical Co., Ltd. (690573662)

**Registrant** - The Mentholatum Company (002105757)

**Establishment**

<b>Name</b>	<b>Address</b>	<b>ID/FEI</b>	<b>Business Operations</b>
The Mentholatum Company		002105757	label(66613-8459)

**Establishment**

<b>Name</b>	<b>Address</b>	<b>ID/FEI</b>	<b>Business Operations</b>
Rohto Pharmaceutical Co., Ltd.		696604024	manufacture(66613-8459)

Revised: 12/2024

Rohto Pharmaceutical Co., Ltd.