# ROHTO COOL RELIEF- naphazoline hydrochloride, polysorbate 80 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.

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#### **Drug Facts**

#### **Active ingredients**

Naphazoline hydrochloride 0.012%

Polysorbate 80 0.2%

#### **Purpose**

Naphazoline hydrochloride - Redness reliever

Polysorbate 80 - Lubricant

#### Uses

- relieves redness of the eye due to minor eye irritations
- for use as a lubricant to prevent further irritation or to relieve dryness of the eye
- for protection against further irritation

#### **Warnings**

#### For external use only

#### Ask a doctor before use if you have

narrow angle glaucoma

#### 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
- overuse may cause more eye redness
- pupils may become enlarged temporarily
- 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) up to 4 times daily
- tightly snap on cap to seal

#### Other information

• do not store above 25°C (77°F)

#### **Inactive ingredients**

alcohol (0.1%), benzalkonium chloride, boric acid, chlorobutanol, edetate disodium, menthol, purified water, sodium borate

#### **Questions?**

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

#### Package/Label Principal Display Panel



#### **ROHTO COOL RELIEF**

naphazoline hydrochloride, polysorbate 80 liquid

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:10742-8141	
Route of Administration	OPHTHALMIC			

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
NAPHAZOLINE HYDROCHLORIDE (UNII: MZ 1131787D) (NAPHAZOLINE - UNII: H231GF11BV)	NAPHAZ OLINE HYDROCHLORIDE	0.12 mg in 1 mL	
POLYSORBATE 80 (UNII: 60ZP39ZG8H) (POLYSORBATE 80 - UNII:60ZP39ZG8H)	POLYSORBATE 80	2 mg in 1 mL	

Inactive Ingredients			
Ingredient Name	Strength		
ALCOHOL (UNII: 3K9958V90M)	1 mg in 1 mL		
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)			
BORIC ACID (UNII: R57ZHV85D4)			
CHLOROBUTANOL (UNII: HM4YQM8WRC)			
EDETATE DISODIUM (UNII: 7FLD91C86K)			
MENTHOL, UNSPECIFIED FORM (UNII: L7T10EIP3A)			
WATER (UNII: 059QF0KO0R)			
SODIUM BORATE (UNII: 91MBZ8H3QO)			

P	Packaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:10742- 8141-1	1 in 1 CARTON	06/21/2021	
1		13 mL in 1 BOTTLE, WTH APPLICATOR; 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	06/21/2021		

## Labeler - The Mentholatum Company (002105757)

### Registrant - The Mentholatum Company (002105757)

Establishment				
Name	Address	ID/FEI	<b>Business Operations</b>	
Rohto-Mentholatum (Vietnam) Co. Ltd.		555347535	manufacture(10742-8141)	

Revised: 2/2023 The Mentholatum Company