

DROPTICS PERFORMANCE EYE DROPS- naphazoline hydrochloride, propylene glycol liquid
Droptics LLC

Active Ingredients

Naphazoline Hydrochloride 0.03%
Propylene Glycol: 1.0%

Purpose

Redness Reliever
Lubricant

Uses

Temporarily relieves burning and irritation due to dryness of the eye
Relieves redness of the eye due to minor eye irritations

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 each 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 use

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 screw on cap to seal after use

Other information

Store at 20 - 25°C (68 - 77°F)

Inactive ingredients

benzalkonium chloride, boric acid, borneol, caffeine, camphor, menthol, potassium l-aspartate, purified water, taurine

Product label



DROPTICS PERFORMANCE EYE DROPS

naphazoline hydrochloride, propylene glycol liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:85211-001
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
NAPHAZOLINE HYDROCHLORIDE (UNII: MZ1131787D) (NAPHAZOLINE - UNII:H231GF11BV)	NAPHAZOLINE HYDROCHLORIDE	0.03 g in 100 mL
PROPYLENE GLYCOL (UNII: 6DC9Q167V3) (PROPYLENE GLYCOL - UNII:6DC9Q167V3)	PROPYLENE GLYCOL	1 g in 100 mL

Inactive Ingredients	
Ingredient Name	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
BORIC ACID (UNII: R57ZHV85D4)	
BORNEOL (UNII: M89NIB437X)	
CAFFEINE (UNII: 3G6A5W338E)	
CAMPHOR (SYNTHETIC) (UNII: 5TJD82A1ET)	
MENTHOL (UNII: L7T10EIP3A)	
POTASSIUM ASPARTATE (UNII: OC4598NZEQ)	
WATER (UNII: 059QF0KO0R)	
TAURINE (UNII: 1EQV5MLY3D)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:85211-001-01	1 in 1 CARTON	03/22/2025	
1		5 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	03/22/2025	

Labeler - Droptics LLC (119445348)

Revised: 3/2025

Droptics LLC