ADVANCED EYE RELIEF/ REDNESS MAXIMUM RELIEF- naphazoline hydrochloride and hypromellose solution/ drops
Bausch & Lomb Incorporated

Drug Facts

Active ingredients

Hypromellose (0.5%)
Naphazoline hydrochloride (0.03%)

Purpose

Lubricant
Redness reliever

Uses

- temporary relief of redness and discomfort due to:
  - minor eye irritations
  - exposure to wind or sun
  - dryness of the eye
- prevents further irritation

Warnings

For external use only

Do not use

- if solution changes color or becomes cloudy

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
- do not overuse as it may produce increased redness of the eye
- pupils may become enlarged temporarily
- remove contact lenses before using
- replace cap after use

Stop use and ask a doctor if

- you experience eye pain, changes in vision, continued redness or irritation of the eye
- 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

- instill 1 to 2 drops in the affected eye(s) up to four times daily

Other information

- store at 15–25 °C (59–77 °F)
- keep tightly closed
- use before expiration date marked on the carton and bottle

Inactive ingredients

benzalkonium chloride, boric acid, edetate disodium, sodium borate, sodium chloride, water for injection. Hydrochloric acid and/or sodium hydroxide may be used to adjust pH.

Questions or comments?

[phone icon] **Call: 1-800-553-5340**

Package/Label Principal Display Panel

BAUSCH + LOMB
ADVANCED
Eye Relief ®
Maximum

Redness
REDNESS RELIEVER/
LUBRICANT EYE DROPS

- Maximum strength redness reliever
- Moisturizes and comforts

STERILE 0.5 FL OZ (15 mL)

3843903
AB55411A

ADVANCED EYE RELIEF/ REDNESS MAXIMUM RELIEF

naphazoline hydrochloride and hypromellose solution/ drops

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:24208-450
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO) (HYPROMELLOSE, UNSPECIFIED - UNII:3NXW29V3WO)	HYPROMELLOSE, UNSPECIFIED	5 mg in 1 mL
NAPHAZOLINE HYDROCHLORIDE (UNII: MZ1131787D) (NAPHAZOLINE - UNII:H231GF11BV)	NAPHAZOLINE HYDROCHLORIDE	0.3 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
BORIC ACID (UNII: R57ZHV85D4)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
WATER (UNII: 059QF0KO0R)	
SODIUM BORATE (UNII: 91MBZ8H3QO)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
HYDROCHLORIC ACID (UNII: QTT17582CB)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:24208-450-15	1 in 1 CARTON	09/02/2010	
1		15 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product		
2	NDC:24208-450-90	2 in 1 CARTON	07/31/2022	07/31/2022

2		15 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	09/02/2010	

Labeler - Bausch & Lomb Incorporated (196603781)

Establishment

Name	Address	ID/FEI	Business Operations
Bausch & Lomb Incorporated		079587625	manufacture(24208-450)

Revised: 9/2024 Bausch & Lomb Incorporated