HEB EYE DROPS- glycerin, naphazoline hydrochloride liquid
H E B

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.

HEB Eye Drops

Active ingredients

Glycerin 0.5%

Naphazoline hydrochloride 0.03%

Purpose

Lubricant

Redness reliever

Uses

- for the relief of redness of the eye due to minor eye irritations.
- for the temporary relief of burning and irritation due to the dryness of the eye.
- for the use as a protectant against further irritation or dryness of the eye

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

- To avoid contamination, do not touch tip of container to any surface.
- Replace cap after using.
- Overuse may produce increased redness of the eye
- Pupils may become enlarged temporarily.

Stop use and ask a doctor if

- you experience eye pain
- you experience changes in vision
- you experience 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 or 2 drops in the affected eye(s) up to 4 times daily.

Other information

Store at room temperature.

Inactive ingredients

benzalkonium chloride, boric acid, disodium edetate hydrate, sodium borate, water for injection

Package label

package label

HEB EYE DROPS

glycerin, naphazoline hydrochloride liquid

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:37808-560
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
NAPHAZOLINE HYDROCHLORIDE (UNII: MZ1131787D) (NAPHAZOLINE - UNII:H231GF11BV)	NAPHAZOLINE HYDROCHLORIDE	0.12 mg in 15 mL
GLYCERIN (UNII: PDC6A3C0OX) (GLYCERIN - UNII:PDC6A3C0OX)	GLYCERIN	2.5 mg in 15 mL

Inactive Ingredients

Ingredient Name	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
BORIC ACID (UNII: R57ZHV85D4)	
SODIUM BORATE (UNII: 91MBZ8H3QO)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:37808-560-02	2 in 1 BOX	12/11/2020	
1		15 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product		
2	NDC:37808-560-01	1 in 1 BOX	12/11/2020	
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 final	part349	12/11/2020	

Labeler - H E B (007924756)

Revised: 1/2023

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