VISINE ORIGINAL REDNESS RELIEF- tetrahydrozoline hydrochloride solution/ drops
Johnson & Johnson Consumer Inc.

Visine® Original Redness Relief Eye Drops

Drug Facts

Active ingredients

Tetrahydrozoline HCl 0.05%

Purpose

Redness reliever

Use

- for the relief of 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

- pupils may become enlarged temporarily
- overuse may cause more eye redness
- remove contact lenses before using
- do not use if this solution changes color or becomes cloudy
- do not touch tip of container to any surface to avoid contamination
- replace cap after each use

Stop use and ask a doctor if

- you feel eye pain
- changes in vision occur
- redness or irritation of the eye lasts
- condition worsens or lasts more than 72 hours

If pregnant or breast-feeding, ask a health professional before use.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions

- put 1 to 2 drops in the affected eye(s) up to 4 times daily
- children under 6 years of age: ask a doctor

Other information

- store at 15° to 25°C (59° to 77°F)

Inactive ingredients

benzalkonium chloride, boric acid, edetate disodium, purified water, sodium borate, sodium chloride

Questions?

call toll-free 888-734-7648 or 215-273-8755 (collect)

Dist: Johnson & Johnson Healthcare Products Division of McNEIL-PPC, Inc., Skillman, NJ 08558 USA

PRINCIPAL DISPLAY PANEL - 15 mL Bottle Carton

Sterile
Visine®

ORIGINAL
Redness Relief
TETRAHYDROZOLINE HCl
REDNESS RELIEVER EYE DROPS

Gets the
Red Out®

Fast-acting
formula

1/2 FL OZ (15mL)

Visine® Original Redness Relief Eye Drops

VISINE ORIGINAL REDNESS RELIEF
tetrahydrozoline hydrochloride solution/ drops

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:42002-203
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
Tetrahydrozoline Hydrochloride (UNII: 0YZT43HS7D) (Tetrahydrozoline - UNII:S9U025Y077)	Tetrahydrozoline Hydrochloride	0.5 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)

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:42002-203-02	1 in 1 CARTON	10/01/2009	09/30/2022
1		15 mL in 1 BOTTLE, DROPPER; Type 4: Device Coated/Impregnated/Otherwise Combined with Drug		
2	NDC:42002-203-35	1 in 1 CARTON	10/01/2009	08/31/2016
2		19 mL in 1 BOTTLE, DROPPER; Type 4: Device Coated/Impregnated/Otherwise Combined with Drug		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH FINAL	part349	10/01/2009	09/30/2022

Labeler - Johnson & Johnson Consumer Inc. (002347102)

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