

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

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.

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

tetrahydrozoline hydrochloride solution/ drops

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:420 02-203
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
Tetrahydrozoline Hydrochloride (UNII: 0 YZT43HS7D) (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: 059QF0K00R)	
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)

Revised: 9/2020

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