

**VISINE RED EYE TOTAL COMFORT MULTI-SYMPTOM- tetrahydrozoline hydrochloride, polyethylene glycol 400, and zinc sulfate, unspecified form 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.*

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**VISINE® Red Eye Total Comfort Multi-Symptom**

***Drug Facts***

<b><i>Active ingredients</i></b>	<b><i>Purpose</i></b>
Polyethylene glycol 400 1%	Lubricant
Tetrahydrozoline HCl 0.05%	Redness reliever
Zinc sulfate 0.25%	Astringent

**Uses**

- for the temporary relief of discomfort and redness of the eye due to minor eye irritations
- relieves dryness of the eye
- for the temporary relief of burning and irritation due to exposure to wind or sun
- for protection against further irritation

**Warnings**

**For external use only**

**Ask a doctor before use if you have** narrow angle glaucoma.

**When using this product**

- some users may experience a brief tingling sensation
- 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**

- adults and children 6 years of age and over: put 1 or 2 drops in the affected eye(s) up to 4 times a day
- children under 6 years of age: consult a doctor
- children under 2 years of age: do not use

### **Other information**

Store at 20° to 25°C (68° to 77°F)

### **Inactive ingredients**

benzalkonium chloride, boric acid, edetate disodium, glycerin, hypromellose, purified water, sodium chloride, sodium citrate

### **Questions?**

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

Distributed by: **JOHNSON & JOHNSON CONSUMER INC.** Skillman, NJ 08558

### **PRINCIPAL DISPLAY PANEL - 15 mL Bottle Carton**

**UP TO 10 HRS COMFORT**

**VISINE®**

**RED EYE**

**TOTAL COMFORT**

**MULTI-SYMPTOM**

**ASTRINGENT /LUBRICANT**

**REDNESS RELIEVER EYE DROPS**

**ALL-IN ONE RELIEF**

From Red, Burning, Watery,

Itchy, Dry, Gritty, Irritated Eyes

Protects Against

Further Irritation

**STERILE**

1/2 FL OZ (15 mL)



## VISINE RED EYE TOTAL COMFORT MULTI-SYMP TOM

tetrahydrozoline hydrochloride, polyethylene glycol 400, and zinc sulfate, unspecified form solution/ drops

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:69968-0360
<b>Route of Administration</b>	OPHTHALMIC		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>TETRAHYDROZOLINE HYDROCHLORIDE</b> (UNII: 0 YZT43HS7D) (TETRAHYDROZOLINE - UNII:S9U025Y077)	TETRAHYDROZOLINE HYDROCHLORIDE	0.5 mg in 1 mL
<b>POLYETHYLENE GLYCOL 400</b> (UNII: B697894SGQ) (POLYETHYLENE GLYCOL	POLYETHYLENE GLYCOL 400	10 mg

400 - UNII:B697894SGQ)	POLYETHYLENE GLYCOL 400	in 1 mL
ZINC SULFATE, UNSPECIFIED FORM (UNII: 89DS0H96TB) (ZINC CATION - UNII:13S1S8SF37)	ZINC SULFATE, UNSPECIFIED FORM	2.5 mg in 1 mL

### Inactive Ingredients

Ingredient Name	Strength
Benzalkonium Chloride (UNII: F5UM2KM3W7)	
Boric Acid (UNII: R57ZHV85D4)	
Edetate Disodium (UNII: 7FLD91C86K)	
Glycerin (UNII: PDC6A3C0OX)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
WATER (UNII: 059QF0KO0R)	
Sodium Chloride (UNII: 451W47IQ8X)	
SODIUM CITRATE, UNSPECIFIED FORM (UNII: 1Q73Q2JULR)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69968-0360-1	1 in 1 CARTON	04/27/2020	
1		15 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	04/27/2020	

**Labeler** - Johnson & Johnson Consumer Inc. (002347102)

Revised: 2/2020

Johnson & Johnson Consumer Inc.