

**VISINE TOTALITY MULTI SYMPTOM RELIEF- glycerin, hypromelloses, polyethylene glycol 400, tetrahydrozoline hydrochloride, and zinc sulfate liquid**  
**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® Totality® Multi-Symptom Relief**

**Drug Facts**

<b>Active ingredients</b>	<b>Purpose</b>
Glycerin 0.2%	Lubricant
Hypromellose 0.36%	Lubricant
Polyethylene glycol 400 1%	Lubricant
Tetrahydrozoline HCl 0.05%	Redness reliever
Zinc Sulfate	Astringent

**Uses**

- for the 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**

- 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 a daily

- children under 6 years of age: ask a doctor

### **Other information**

- some users may experience a brief tingling sensation
- store at 20° to 25°C (68° to 77°F)

### **Inactive ingredients**

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

### **Questions?**

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

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

*Sterile*

**Visine®**

**TOTALITY®**

**Multi-Symptom  
Relief**

**LUBRICANT / ASTRINGENT /  
REDNESS RELIEVER  
EYE DROPS**

UP

TO

10

HOURS

COMFORT

HYDROBLEND®

Relieves

red, burning,

watery, itchy,

gritty, dry,

irritated eyes

with HYDROBLEND®

moisturizers

for up to 10 hours

of comfort

1/2 FL OZ (15mL)



## VISINE TOTALITY MULTI SYMPTOM RELIEF

glycerin, hypromelloses, polyethylene glycol 400, tetrahydrozoline hydrochloride, and zinc sulfate liquid

### Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:42002-209
Route of Administration	OPHTHALMIC		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>Glycerin</b> (UNII: PDC6A3C0OX) (Glycerin - UNII:PDC6A3C0OX)	Glycerin	2.5 mg in 1 mL
<b>HYPROMELLOSE, UNSPECIFIED</b> (UNII: 3NXW29V3WO) (HYPROMELLOSE, UNSPECIFIED - UNII:3NXW29V3WO)	HYPROMELLOSE, UNSPECIFIED	3.6 mg in 1 mL
<b>Polyethylene Glycol 400</b> (UNII: B697894SGQ) (Polyethylene Glycol 400 -	Polyethylene Glycol 400	11.28 mg

UNII:B697894SGQ)	Polyethylene Glycol 400	in 1 mL
<b>Tetrahydrozoline Hydrochloride</b> (UNII: 0YZT43HS7D) (Tetrahydrozoline - UNII:S9U025Y077)	Tetrahydrozoline Hydrochloride	0.5 mg in 1 mL
<b>Zinc Sulfate, Unspecified Form</b> (UNII: 89DS0H96TB) (Zinc Cation - UNII:13S1S8SF37)	Zinc Sulfate, Unspecified Form	2.5 mg in 1 mL

### Inactive Ingredients

Ingredient Name	Strength
<b>Benzalkonium Chloride</b> (UNII: F5UM2KM3W7)	
<b>Boric Acid</b> (UNII: R57ZHV85D4)	
<b>Edetate Disodium</b> (UNII: 7FLD91C86K)	
<b>Water</b> (UNII: 059QF0K00R)	
<b>Sodium Chloride</b> (UNII: 451W47IQ8X)	
<b>Sodium Citrate, Unspecified Form</b> (UNII: 1Q73Q2JULR)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:42002-209-05	1 in 1 CARTON	08/23/2011	08/31/2021
1		15 mL in 1 BOTTLE; 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	08/23/2011	08/31/2021

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

Revised: 1/2020

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