

**MEIJER MULTI-SYMPTOM RELIEF- polyethylene glycol, tetrahydrozoline hydrochloride, zinc sulfate solution/ drops**  
**K.C. Pharmaceuticals, 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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**Meijer Multi-Symptom Relief**

**DRUG FACTS**

**Active ingredients**

Polyethylene glycol 400 .....1%  
Tetrahydrozoline HCl.....0.05%  
Zinc sulfate.....0.25%

**Purposes**

Polyethylene glycol 400 .....Lubricant  
Tetrahydrozoline HCl.....Redness reliever  
Zinc sulfate.....Astringent

**Uses**

- relieves dryness of the eye
- for temporary relief of discomfort and redness of the eye due to minor eye irritations
- for 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 touch tip of container to any surface to avoid contamination
- replace cap after using
- do not use if this solution changes color or becomes cloudy

**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 (1-800-222-1222) right away.

**Directions**

- Instill 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**

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

**Inactive ingredients**

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

**Questions or comments?**

**Call 1-888-527-4276**

**meijer** multi-symptom relief  
Lubricant/Astringent/Redness Reliever Eye Drops

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\*This product is not manufactured or distributed by Johnson & Johnson, Healthcare Products, owner of the registered trademark Visine®.

**meijer** NDC 79481-0250-1  
Compare to Visine® Red Eye Total Comfort Multi-Symptom active ingredients\*

**multi-symptom relief**  
Polyethylene glycol 400 1%  
Tetrahydrozoline HCl 0.05%  
Zinc sulfate 0.25%

**Lubricant/Astringent/Redness Reliever Eye Drops**

**STERILE**

Relieves red, burning, watery, itchy, gritty, dry, irritated eyes

0.5 FL OZ (15 mL)

**OUR QUALITY GUARANTEE**  
The Meijer Family  
WWW.MEIJER.COM/SATISFACTION

**PAPER BOX** **PLASTIC BOTTLE**

PHD 5159810  
13733 20139 1  
CEDVT0051MEEC

**MEIJER MULTI-SYMPATOM RELIEF**

polyethylene glycol, tetrahydrozoline hydrochloride, zinc sulfate solution/ drops

## Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:55651-250
<b>Route of Administration</b>	OPHTHALMIC		

## Active Ingredient/Active Moiety

<b>Ingredient Name</b>	<b>Basis of Strength</b>	<b>Strength</b>
<b>TETRAHYDROZOLINE HYDROCHLORIDE</b> (UNII: 0YZT43HS7D) (TETRAHYDROZOLINE - UNII:S9U025Y077)	TETRAHYDROZOLINE HYDROCHLORIDE	0.05 g in 100 mL
<b>POLYETHYLENE GLYCOL 400</b> (UNII: B697894SGQ) (POLYETHYLENE GLYCOL 400 - UNII:B697894SGQ)	POLYETHYLENE GLYCOL 400	1 g in 100 mL
<b>ZINC SULFATE</b> (UNII: 89DS0H96TB) (ZINC CATION - UNII:13S1S8SF37)	ZINC SULFATE	0.25 g in 100 mL

## Inactive Ingredients

<b>Ingredient Name</b>	<b>Strength</b>
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>SODIUM CITRATE</b> (UNII: 1Q73Q2JULR)	
<b>BORIC ACID</b> (UNII: R57ZHV85D4)	
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	
<b>BENZALKONIUM CHLORIDE</b> (UNII: F5UM2KM3W7)	
<b>EDETATE DISODIUM</b> (UNII: 7FLD91C86K)	
<b>HYPROMELLOSES</b> (UNII: 3NXW29V3WO)	
<b>SODIUM CHLORIDE</b> (UNII: 451W47IQ8X)	

## Packaging

<b>#</b>	<b>Item Code</b>	<b>Package Description</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
1	NDC:55651-250-01	1 in 1 CARTON	01/09/2023	
1		15 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product		

## Marketing Information

<b>Marketing Category</b>	<b>Application Number or Monograph Citation</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
OTC monograph final	part349	01/09/2023	

**Labeler** - K.C. Pharmaceuticals, Inc. (174450460)

## Establishment

<b>Name</b>	<b>Address</b>	<b>ID/FEI</b>	<b>Business Operations</b>
K.C. Pharmaceuticals, Inc.		174450460	manufacture(55651-250) , pack(55651-250) , label(55651-250)

