

**EQUATE REDNESS RELIEVER- tetrahydrozoline hydrochloride solution/ drops  
WALMART 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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**Equate Redness Relief eye drops - tetrahydrozoline HCl**

**Drug Facts**

***Active ingredient***

Tetrahydrozoline Hydrochloride 0.05%

***Purpose***

Redness reliever

***Uses***

- relieves redness of the eye due to minor eye irritations

***Warnings***

**For external use only**

**Do not use**

- if solution changes color or becomes cloudy

**Ask a doctor before use if you have**

- narrow angle glaucoma

**When using this product**

- to avoid contamination, do not touch tip of container to any surface
- replace cap after using
- overuse may produce increased redness of the eye
- pupils may become enlarged temporarily

**Stop use and ask a doctor if**

- you experience eye pain
- you experience changes in vision
- you experience continued redness or irritation of the eye
- the condition worsens or persists for more than 72 hours

**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 four times daily

***Other information***

- store at room temperature
- remove contact lenses before using

***Inactive ingredients***

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

***Questions?***

1-888-287-1915

**Principal Display Panel**

**Equate<sup>®</sup>  
Original Redness Reliever  
Tetrahydrozoline HCl  
Sterile Eye Drops  
1 FL OZ (30 mL)**



**Principal Display Panel**

**Equate®**  
**Original Redness**  
**Reliever**  
**Tetrahydrozoline HCl**  
**Sterile Eye Drops**  
**0.5 FL OZ (15 mL)**



## EQUATE REDNESS RELIEVER

tetrahydrozoline hydrochloride solution/ drops

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:79903-010
<b>Route of Administration</b>	OPHTHALMIC		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>TETRAHYDROZOLINE HYDROCHLORIDE</b> (UNII: 0YZT43HS7D) (TETRAHYDROZOLINE - UNII:S9U025Y077)	TETRAHYDROZOLINE HYDROCHLORIDE	0.5 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: 059QF0KO0R)	
<b>SODIUM BORATE</b> (UNII: 91MBZ8H3QO)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:79903-010-15	1 in 1 CARTON	04/26/2013	
1		15 mL in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:79903-010-30	1 in 1 CARTON	04/26/2013	
2		30 mL in 1 BOTTLE; Type 0: Not a Combination Product		

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part349	04/26/2013	

**Labeler** - WALMART INC. (051957769)

Revised: 1/2022

WALMART INC.