

ORIGINAL REDNESS RELIEVER- tetrahydrozoline hcl solution/ drops
Walmart

Equate Original Redness Reliever

Purpose

Redness Reliever

Active ingredient

TetrahydrozolineHCl0.05%

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center (1-800-222-1222) right away.

Inactive ingredients

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

Directions

instill 1 to 2 drops in the affected eye(s) up to 4times daily

■ relieves 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

■ to avoid contamination, do not touch tip of container to any surface. Replace cap after using.

■ if solution changes color or becomes cloudy, do not use

■ overuse may produce increased redness of the eye

■ remove contact lenses before using

Stop use and ask a doctor if you experience

■ eye pain

■ changes in vision

■ continued redness or irritation of the eye, or if the condition worsens or persists for more

than 72 hours

Hpregnant orbreast-feeding,askahealth professional before use.



ORIGINAL REDNESS RELIEVER

tetrahydrozoline hcl solution/ drops

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC: 79903-363
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
TETRAHYDROZOLINE HYDROCHLORIDE (UNII: 0YZT43HS7D) (TETRAHYDROZOLINE - UNII:S9U025Y077)	TETRAHYDROZOLINE HYDROCHLORIDE	0.05 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
SODIUM BORATE (UNII: 91MBZ8H3QO)	
WATER (UNII: 059QF0KO0R)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
BORIC ACID (UNII: R57ZHV85D4)	
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:79903-363-01	1 in 1 BOX	12/26/2025	
1		15 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 Drug	M018	12/26/2025	

Labeler - Walmart (051957769)

Revised: 12/2025

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