MEIJER SEASONAL ITCHING AND REDNESS RELIEF- tetrahydrozoline hci, zinc sulfate solution/ drops Meijer Distribution, Inc.

Meijer Seasonal Itching & Redness Relief (PLD)

Active ingredients

Tetrahydrozoline HCI 0.05%

Zinc sulfate 0.25%

Purposes

Redness reliever

Astringent

Use

for temporary relief of discomfort and 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 lens 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

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

Other information

store at 15°-30°C (59°-86°F)

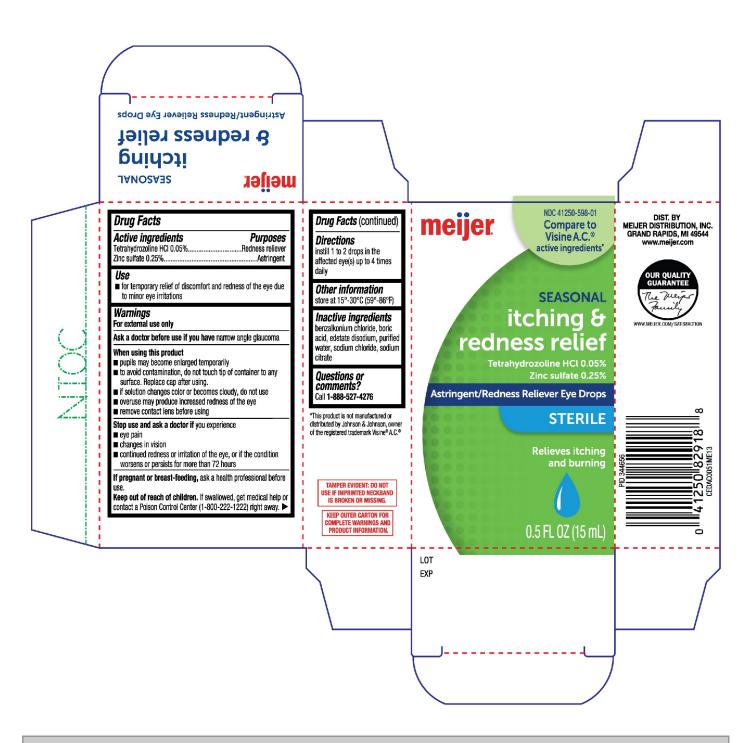
Inactive ingredients

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

Questions or comments?

Call 1-888-527-4276

Meijer Seasonal Itching & Redness Relief 15mL



MEIJER SEASONAL ITCHING AND REDNESS RELIEF

tetrahydrozoline hci, zinc sulfate solution/ drops

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Product Type HUMAN OTC DRUG Item Code (Source) NDC:41250-598

Route of Administration OPHTHALMIC

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

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Inactive Ingredients		
Ingredient Name	Strength	
EDETATE DISODIUM (UNII: 7FLD91C86K)		
SODIUM CHLORIDE (UNII: 451W47IQ8X)		
BORIC ACID (UNII: R57ZHV85D4)		
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)		
WATER (UNII: 059QF0KO0R)		
SODIUM CITRATE (UNII: 1Q73Q2JULR)		

ı	Packaging			
	# Item Code	Package Description	Marketing Start Date	Marketing End Date
	1 NDC:41250- 598-01	1 in 1 BOX	03/10/2020	
	1	15 mL in 1 BOTTLE, DROPPER; 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	03/10/2020	

Labeler - Meijer Distribution, Inc. (006959555)

Registrant - KC Pharmaceuticals, Inc. (174450460)

Establishment			
Name	Address	ID/FEI	Business Operations
KC Pharmaceuticals, Inc.		174450460	manufacture(41250-598) , pack(41250-598) , label(41250-598)

Revised: 12/2023 Meijer Distribution, Inc.