

SUNMARK EYE DROPS ORIGINAL FORMULA- tetrahydrozoline hcl solution/ drops

Strategic Sourcing Services LLC

Sunmark Eye Drops Original Formula (PLD)

Active Ingredient

Tetrahydrozoline HCL 0.05%

Purpose

Redness reliever

Uses

- 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 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 right away

Directions

Instill 1 or 2 drops in the affected eye(s) up to four times daily

Other information

Store at 15 -30°C (59°-86°F)

Inactive ingredients

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



FAST ACTING

Original Formula
sterile
eye drops
sunmark™

Drug Facts

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Drug Facts (continued)

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sunmark™

NDC 49348-037-29

sterile
eye drops

Original Formula

Fast acting
Redness reliever

FAST ACTING

1/2 FLOZ (15 mL)

■ Tamper Evident. Do not use this product if imprinted neckband is missing or broken.

■ RETAIN THIS CARTON FOR FUTURE REFERENCE

McKesson
Empowering HealthCare

Another Quality Product
Distributed By McKesson
One Post Street
San Francisco, CA 94104
Money Back Guarantee
Please visit us at
www.sunmarkbrand.com

CEDRG0051SM0



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SUNMARK EYE DROPS ORIGINAL FORMULA

tetrahydrozoline hcl solution/ drops

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:49348-037
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
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
BORIC ACID (UNII: R57ZHV85D4)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
WATER (UNII: 059QF0KO0R)	
SODIUM BORATE (UNII: 91MBZ8H3QO)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:49348-037-29	1 in 1 CARTON	03/01/2007	
1		15 mL in 1 BOTTLE, PLASTIC; 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	03/01/2007	

Labeler - Strategic Sourcing Services LLC (116956644)

Registrant - KC Pharmaceuticals, Inc. (174450460)

Establishment

Name	Address	ID/FEI	Business Operations
KC Pharmaceuticals, Inc.		174450460	manufacture(49348-037) , pack(49348-037) , label(49348-037)