

**MEDICS CHOICE EYE DROPS ADVANCED- tetrahydrozoline hydrochloride, polyethylene glycol 400, dextran 70, povidone solution/ drops
KC Pharmaceuticals, Inc.**

Medic's Choice Eye Drops Advanced

Active ingredients

Dextran 70 0.1%

Polyethylene glycol 400 1%

Povidone 1%

Tetrahydrozoline HCl 0.05%

Purposes

Dextran 70.....Lubricant

Polyethylene glycol 400....Lubricant

Povidone.....Lubricant

Tetrahydrozoline HCl.....Redness reliever

Uses

- relieves redness of the eye due to minor eye irritations
- as a lubricant to prevent further irritation or to relieve dryness of the eye

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 borate, and sodium chloride

Medic's Choice EYE DROPS ADVANCED

Drug Facts

Active ingredients	Purposes
Dextran 70 0.1%.....	Lubricant
Polyethylene glycol 400 1%.....	Lubricant
Povidone 1%.....	Lubricant
Tetrahydrozoline HCl 0.05%.....	Redness reliever

Uses

- relieves redness of the eye due to minor eye irritations
- as a lubricant to prevent further irritation or to relieve dryness of the eye

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.

Drug Facts (continued)

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 borate, and sodium chloride

*This product is not manufactured or distributed by Johnson & Johnson, Healthcare Products, distributor of Visine® Advanced Relief.

TAMPER EVIDENT: DO NOT USE IF IMPRINTED NECKBAND IS BROKEN OR MISSING.

KEEP OUTER CARTON FOR COMPLETE WARNINGS AND PRODUCT INFORMATION.

Compare to the active ingredients in Visine® Advanced Relief*

Medic's Choice. EYE DROPS ADVANCED

Lubricant Redness Reliever

- Cools
- Soothes
- Refreshes

Sterile 0.5 fl. oz. (15 mL)

LOT EXP

NDC 55651-025-01

KC Pharmaceuticals, Inc. Pomona, CA 91768 Questions? Call 1-888-527-4276 Made in USA

0 95072 01251 8 CEDAR0051MC6

MEDICS CHOICE EYE DROPS ADVANCED

tetrahydrozoline hydrochloride, polyethylene glycol 400, dextran 70, povidone solution/ drops

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:55651-025
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
POVIDONE (UNII: FZ989GH94E) (POVIDONE - UNII:FZ989GH94E)	POVIDONE	10 mg in 1 mL
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ) (POLYETHYLENE GLYCOL 400 - UNII:B697894SGQ)	POLYETHYLENE GLYCOL 400	10 mg in 1 mL
TETRAHYDROZOLINE HYDROCHLORIDE (UNII: 0YZT43HS7D) (TETRAHYDROZOLINE - UNII:S9U025Y077)	TETRAHYDROZOLINE HYDROCHLORIDE	0.5 mg in 1 mL
DEXTRAN 70 (UNII: 7SA290YK68) (DEXTRAN 70 - UNII:7SA290YK68)	DEXTRAN 70	1 mg in 1 mL

Inactive Ingredients

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

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:55651-025-01	1 in 1 CARTON	12/09/2003	
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	12/09/2003	

Labeler - KC Pharmaceuticals, Inc. (174450460)

Establishment

Name	Address	ID/FEI	Business Operations
KC Pharmaceuticals, Inc.		174450460	manufacture(55651-025) , pack(55651-025) , label(55651-025)

Revised: 1/2024

KC Pharmaceuticals, Inc.