MULTI-ACTION RELIEF- polyvinyl alcohol, povidone and tetrahydrozoline hydrochloride solution/ drops
Altaire Pharmaceuticals Inc.

Multi-Action Relief Drops

Multi-Action Relief Drops15mLNDC 59390-156-13

Drug Facts

Active ingredients

Polyvinyl alcohol 0.5%

Povidone 0.6%

Tetrahydrozoline hydrochloride 0.05%

Purpose

Lubricant

Lubricant

Redness Reliever

Uses

- for the temporary relief of burning and irritation due to dryness of the eye
- for use as a protectant against further irritation or to relieve dryness of the eye
- relieves redness of the eye due to minor eye irritations

Warnings

Forusein the eye(s)only.

Do not useif

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 ofcontainer 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
- symptoms last for more than 72 hours

Keepoutofreach of children.

If swallowed, get medical helporcontact a Poison Control Center right away.

Directions

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

OtherInformation

- store at room temperature.
- remove contact lenses before using
- tamper evident: Do not use if imprinted neckband on bottle is torn, broken or missing.

Inactive ingredients

benzalkonium chloride, dextrose, dibasic sodium
phosphate, edetate disodium, monobasic sodium phosphate, potassium
chloride, purified water, sodium bicarbonate, sodium chloride, sodium citrate.
Hydrochloric acid or sodium hydroxide may be added to adjust pH.

PRINCIPAL DISPLAY PANEL

Multi-Action RELIEF DROPSLUBRICANT/REDNESSRELIEVER EYE DROPSSTERILE0.5 FL OZ (15mL)

MULTI-ACTION RELIEF

polyvinyl alcohol, povidone and tetrahydrozoline hydrochloride solution/ drops

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:59390-156
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
POLYVINYL ALCOHOL, UNSPECIFIED (UNII: 532B59J990) (POLYVINYL ALCOHOL, UNSPECIFIED - UNII:532B59J990)	POLYVINYL ALCOHOL, UNSPECIFIED	5 mg in 1 mL
POVIDONE (UNII: FZ989GH94E) (POVIDONE - UNII:FZ989GH94E)	POVIDONE	6 mg in 1 mL
TETRAHYDROZOLINE HYDROCHLORIDE (UNII: 0YZT43HS7D) (TETRAHYDROZOLINE - UNII:S9U025Y077)	TETRAHYDROZOLINE HYDROCHLORIDE	0.5 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
DEXTROSE (UNII: IY9XDZ35W2)	
SODIUM PHOSPHATE, DIBASIC (UNII: GR686LBA74)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
SODIUM PHOSPHATE, MONOBASIC (UNII: 3980JIH2SW)	
POTASSIUM CHLORIDE (UNII: 660YQ98I10)	
WATER (UNII: 059QF0KO0R)	
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:59390-156-13	1 in 1 CARTON	05/09/2009	
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	05/09/2009	

Labeler - Altaire Pharmaceuticals Inc. (786790378)

Establishment

Name	Address	ID/FEI	Business Operations
Altaire Pharmaceuticals Inc.		786790378	manufacture(59390-156)

Revised: 12/2023 Altaire Pharmaceuticals Inc.