

**STYE- polyvinyl alcohol and povidone solution/ drops
Medtech Products Inc.**

Stye Eye Drops 63029-324

Drug Facts

Active ingredients

Polyvinyl alcohol 0.5%
Povidone 0.6%

Purpose

Lubricant
Lubricant

Uses

For temporary relief of discomfort due to minor irritations including: burning, stinging, itching.

Warnings

for external use only

Do not use

if solution changes color or becomes cloudy

When using this product

- to avoid contamination, do not touch tip to any surface
- replace cap after using

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 or persists for more than 72 hours

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) as needed

Other information

- store at room temperature
- remove contact lenses before using

Inactive ingredients

benzalkonium chloride, dextrose, dibasic sodium phosphate, edetate disodium, monobasic sodium phosphate, potassium chloride, purified water, sodium bicarbonate, sodium chloride, sodium citrate

Questions?

1-800-344-7239

PRINCIPAL DISPLAY PANEL

Stye TM

LUBRICANT EYE DROPS

STERILE 0.5 fl oz (15 mL)



STYE

polyvinyl alcohol and povidone solution/ drops

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63029-324
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

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)	
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63029-324-15	1 in 1 BOTTLE	12/01/2021	
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	12/01/2021	

Labeler - Medtech Products Inc. (122715688)

Revised: 10/2025

Medtech Products Inc.