POVIDONE-IODINE- povidone-iodine solution James Alexander Corporation

POVIDONE IODINE TOPICAL SOLUTION USP

Drug Facts

Active Ingredients (each swab)

10% Povidone Iodine Solution USP, (1% available Iodine)

Purpose

Topical Antiseptic

Uses

To treat minor skin cuts and abrasions.

Warnings

For external use only.

Ask a doctor before use if you have • deep or puncture wounds • serious burns

Stop use and ask a doctor if • redness, irritation, swelling or pain persists or increases

infection occurs.

Keep out of reach of children. If swallowed, get medical help immediately or contact a Poison Control Center right away.

Directions

Reverse cardboard sleeve then crush at dot between thumb and forefinger. Allow solution to saturate tip and apply solution to injury. **Discard after single use.**

Other Information

Store at room temperature away from light. Keep from freezing or excessive heat.

Inactive Ingredients

Citric acid, disodium phosphate, nonoxynol-9, sodium hydroxide, water.

Questions?

Call 1-908-362-9266 Monday through Friday, 9:00am - 5:00pm e.s.t.

DISPENSING SOLUTIONS®

JAMES ALEXANDER CORPORATION

Blairstown, NJ• (908) 362-9266 Product information and MSDS available on-line at: **www.james-alexander.com**

Void of Aqueous

Packaging



Drug Facts



POVIDONE-IODINE

povidone-iodine solution

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:46414-7777

Route of Administration TOPICAL

Active Ingredient/Active Moiety Ingredient Name Basis of Strength

POVIDONE-IODINE (UNII: 85H0HZU99M) (IODINE - UNII:9679TC07X4) IODINE 10 mg in 1 mL

Inactive Ingredients

Ingredient Name Strength

Strength

CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)

SODIUM PHOSPHATE, DIBASIC, ANHYDROUS (UNII: 22ADO53M6F)

NONOXYNOL-9 (UNII: 48Q180SH9T)

SODIUM HYDROXIDE (UNII: 55X04QC32I)

WATER (UNII: 059QF0KO0R)

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#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:46414- 7777-3	10 in 1 CONTAINER	02/14/1976		
1		0.6 mL in 1 VIAL, SINGLE-USE; Type 0: Not a Combination Product			
2	NDC:46414- 7777-2	100 in 1 CONTAINER	02/14/1976		
2		0.6 mL in 1 VIAL, SINGLE-USE; Type 0: Not a Combination Product			

Marketing Information

Marketing Application Number or Monograph Category Citation Date Date

OTC Monograph Drug M003 02/14/1976

Labeler - James Alexander Corporation (040756421)

Registrant - James Alexander Corporation (040756421)

Establishment

Name	Address	ID/FEI	Business Operations

James Alexander Corporation	040756421	manufacture(46414-7777)

Revised: 12/2024 James Alexander Corporation