APLICARE POVIDONE-IODINE SCRUB- povidone-iodine scrub solution
Aplicare Products, LLC

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

0024 Aplicare Povidone-Iodine Scrub

Active ingredient

Povidone-iodine 10%

Purpose

Antiseptic

Use

Antiseptic skin preparation

Warnings

For external use only

Avoid pooling beneath the patient. Prolonged exposure to wet solution may cause skin irritation.

Avoid excessive heat. Store at room temperature.

Ask a doctor before use if injuries are

- deep wounds
- puncture wounds
- serious burns

Stop use and ask a doctor if

- redness, irritation, swelling or pain persists or increases
- infection occurs

Do not use

- if allergic to iodine
- in the eyes

Keep out of reach of children.

In case of accidental ingestion, seek professional assistance or consult a poison control center immediately.

Directions

use sponges to prep desired area

Other information

- for single use only
- not made with natural rubber latex
- for hospital or professiona use only

Questions or comments?

800 633 5463

Inactive ingredients

ammonium nonoxynol-4 sulfate, citric acid, disodium phosphate, nonoxynol-9, sodium hydroxide, water

Manufacturing Information

Manufactured by:

Aplicare Products, LLC

550 Research Pkwy, Meriden, CT 06450 USA

Made in USA with domestic and foreign materials

1-800-633-5463

REF: ORF22025S

RD18APL

Package Label

Tear Here Hold Upright Tear Here

TWO POVIDONE-IODINE SCRUB SPONGE STICKS

ANTISEPTIC

STERILE Solution

Applicator is STERILE if Package is Intact

Two 8-inch Sponge Sticks Saturated with Povidone-Iodine Scrub USP

STERILE SOLUTION

Drug Facts

Active ingredient — **Purpose**
Povidone-Iodine USP 10% ..... Antiseptic

Use antiseptic skin preparation

Warnings
Do not use ■ if allergic to iodine ■ in the eyes

Ask a doctor before use if injuries are
■ deep wounds ■ puncture wounds ■ serious burns

Stop use and ask a doctor if
■ redness, irritation, swelling or pain persists or increases
■ infection occurs

For external use only

Keep out of reach of children. In case of accidental ingestion, seek professional assistance or consult a poison control center immediately.

Avoid pooling beneath the patient. Prolonged exposure to wet solution may cause skin irritation.

Avoid excessive heat. Store at room temperature.

Drug Facts (continued)

Directions use sponge sticks to prep desired area

Other information
■ for single use only
■ not made with natural rubber latex
■ for hospital or professional use only

Inactive ingredients ammonium nonoxynol-4 sulfate, citric acid, disodium phosphate, nonoxynol-9, sodium hydroxide, water

For questions, comments, or to report serious side effects:
1-800-633-5463

Manufactured by:
Aplicare Products, LLC
Meriden, CT 06450
www.medline.com

RD18APL

Reorder No. ORF22025S

(01) 0 03523800 02427

APLICARE POVIDONE-IODINE SCRUB

povidone-iodine scrub solution

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:52380-0024
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
POVIDONE-IODINE (UNII: 85H0HZU99M) (IODINE - UNII:9679TC07X4)	IODINE	10 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
SODIUM PHOSPHATE, DIBASIC (UNII: GR686LBA74)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
NONOXYNOL-9 (UNII: 48Q180SH9T)	
AMMONIUM NONOXYNOL-4 SULFATE (UNII: 9HIA70O4J0)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:52380-0024-2	90 mL in 1 PACKET; Type 0: Not a Combination Product	07/01/1992	02/28/2025
2	NDC:52380-0024-1	120 mL in 1 PACKET; Type 0: Not a Combination Product	07/01/1992	03/31/2025
3	NDC:52380-0024-3	120 mL in 1 PACKET; Type 0: Not a Combination Product	07/01/1992	01/31/2025

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	07/01/1992	03/31/2025

Labeler - Aplicare Products, LLC (081054904)

Registrant - Medline Industries, LP (025460908)

Revised: 12/2022 Aplicare Products, LLC