

POVIDONE-IODINE- povidone-iodine solution
Medline Industries, LP

938 ReadyScrub PVP

Active ingredient

Povidone-Iodine 7.5% w/v

(equivalent to 0.75% available iodine)

Purpose

Antiseptic

Uses

- for preparation of the skin prior to surgery
- helps to reduce bacteria that potentially can cause skin infection
- significantly reduces the number of microorganisms on the hands and forearms prior to surgery or patient care
- single use when used for patient preoperative skin preparation

Warnings

For external use only

Do not use

- in the eyes
- if you are allergic to iodine or any of the other ingredients in the product

When using this product

- prolonged exposure to wet solution may cause irritation or, rarely, severe skin reactions
- in pre-operative prepping, avoid “pooling” beneath the patient

Stop use and ask a doctor

- if irritation and redness develop
- if condition persists for more than 72 hours

Keep out of reach of children.

If swallowed, get medical help or contact a Poison control center right away.

Directions

- apply topically as needed

Other information

- protect from freezing, avoid excessive heat

Inactive ingredients

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

Manufacturing Information

www.medline.com

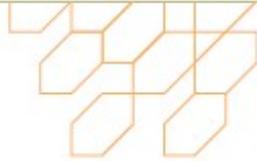
Manufactured in USA by Medline Industries, LP,

Three Lakes Drive, Northfield, IL 60093 USA.

1-800-MEDLINE

REF: MDS093947

Package Label



NDC: 53329-938-06

READYSCRUB™ PVP Povidone-Iodine 7.5% Solution

- Topical Antiseptic
- Surgical Hand Scrub
- Non-sterile
- Not made with Natural Rubber Latex



REF MDS093947

LOT

Exp:



Drug Facts

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1-800-MEDLINE V3 RC25HND

POVIDONE-IODINE

povidone-iodine solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:53329-938
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
POVIDONE-IODINE (UNII: 85H0HZU99M) (IODINE - UNII:9679TC07X4)	IODINE	0.75 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
AMMONIUM NONOXYNOL-4 SULFATE (UNII: 9HIA70O4J0)	
NONOXYNOL-5 (UNII: ED8J5T817W)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
SODIUM PHOSPHATE, DIBASIC, DIHYDRATE (UNII: 94255I6E2T)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:53329-938-06	473 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/06/2017	
2	NDC:53329-938-08	237 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/06/2017	
3	NDC:53329-938-23	946 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/06/2017	
4	NDC:53329-938-25	3785 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/06/2017	
5	NDC:53329-938-04	118 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/06/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	01/01/2007	

Labeler - Medline Industries, LP (025460908)

Registrant - Medline Industries, LP (025460908)

Revised: 10/2025

Medline Industries, LP