

FIRST AID ANTISEPTIC- povidone-iodine solution

Dolgencorp 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.

Povidone-Iodine

050.002 050AE

Active ingredient

Povidone-iodine,
(equivalent to 1% available iodine)

Purpose

First aid antiseptic

Use

first aid to help prevent the risk of infection in

- minor cuts
- scrapes
- burns

Warnings

For external use only

Do not use

- in the eyes or apply over large areas of the body
- longer than 1 week unless directed by a doctor

ask a doctor before use

in case of deep or puncture wounds, animal bites, or serious burns

stop use and ask a doctor if

- condition persists or gets worse

Keep out of reach of children

If swallowed, get medical help or contact a Poison Control Center right away.

Directions

- apply a clean the affected area
- small amount of this product on the area 1 to 3 times daily
- may be covered with a sterile bandage.
- if bandaged, let dry first.

Other information

- store between 15°-30°C (59°-86°F)

Inactive ingredients

C12-13 Pareth-9, citric acid, disodium phosphate, glycerin, sodium hydroxide, water

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Topical Antiseptic

Microbicide

Povidone Iodine Solution, 10%

kills germs in minor burns, cuts and scrapes

4 FL OZ (118 mL)



FIRST AID ANTISEPTIC

povidone-iodine solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:55910-001
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
C12-13 PARETH-9 (UNII: 9BXD858P37)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
SODIUM PHOSPHATE, DIBASIC, ANHYDROUS (UNII: 22ADO53M6F)	
GLYCERIN (UNII: PDC6A3C00X)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:55910-001-26	118 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/11/2012	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	01/11/2012	

Labeler - Dolgencorp LLC (068331990)

Registrant - Vi Jon, LLC (790752542)

Establishment

Name	Address	ID/FEI	Business Operations
Vi Jon, LLC		790752542	manufacture(55910-001)

Establishment

Name	Address	ID/FEI	Business Operations
Vi Jon, LLC		088520668	manufacture(55910-001)

Revised: 4/2022

Dolgencorp LLC