

ZEE POVIDONE IODINE- povidone iodine 10% w/w (equivalent to iodine 1%) swab

Cintas Corporation

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 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.

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, 9:00 am - 5:00 pm e.s.t.

Package Principal Display Panel

10 SWABS

(1/2 ml.)

UNIT NO.

U1-16A

POVIDONE - IODINE SWABS U.S.P

(ANTISEPTIC)

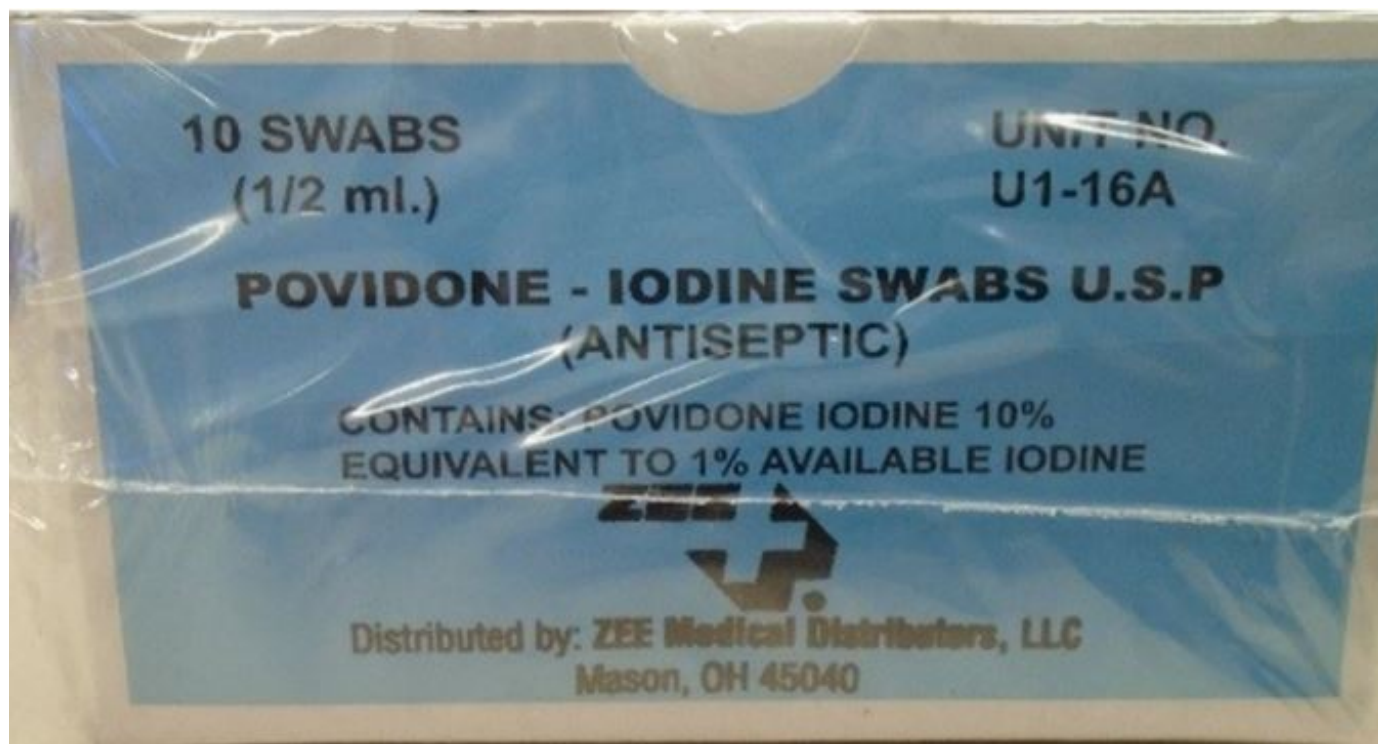
CONTAINS: POVIDONE IODINE 10%

EQUIVALENT TO 1% AVAILABLE IODINE

ZEE®

Distributed by: **ZEE Medical Distributors, LLC**

Mason, OH 45040



ZEE POVIDONE IODINE

povidone iodine 10% w/w (equivalent to iodine 1%) swab

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:42961-023
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
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
SODIUM PHOSPHATE, DIBASIC (UNII: GR686LBA74)	
NONOXYNOL-9 (UNII: 48Q180SH9T)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
WATER (UNII: 059QF0KO0R)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:42961-023-01	10 in 1 BOX	05/12/2022	
1		0.5 mL in 1 VIAL, SINGLE-USE; Type 0: Not a Combination Product		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003	05/12/2022	

Labeler - Cintas Corporation (056481716)

Establishment			
Name	Address	ID/FEI	Business Operations
James Alexander Corporation		040756421	manufacture(42961-023)

Revised: 12/2023

Cintas Corporation