

POVIDONE-IODINE GEL SWABSTICK- povidone iodine gel
Dukal LLC

Dukal Povidone-Iodine Gel Swabstick Non-Sterile

Active Ingredient

Povidone-Iodine USP 10%

Purpose

Antisepetic

Use

Antiseptic Skin Preparation

Warnings

For External Use Only

Do not use

- If allergic to Iodine
- in the eyes

Ask a doctor before use if injuries are

- deep or puncture wounds
- Serious burn

Stop use and ask doctor if

- infection occurs
- redness, irritation or swelling develops

Keep out of reach of children

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

Directions

- apply product to skin as needed
- see product box for additional directions

Other Information

- 1% titratable iodine
- Avoid excessive heat
- Store at room temperature

- For hospital or professional use only
- Not made with natural rubber latex

Inactive Ingredients

alkyl glucoside, anhydrous citric acid, aonoxynol-10, glycerin, hydroxyethyl cellulose, potassium iodine, sodium hydroxide, water

Dukal Povidone-Iodine Gel Swabstick



NDC 65517-0036-1

REF 899

Hold Upright and Tear Here

**POVIDONE-IODINE
GEL SWABSTICK
READY GEL 1X™**



**4" (10.2 cm)
Saturated Swabstick**

Drug Facts

Active ingredient Purpose

Povidone Iodine USP 10% Antiseptic

Use Patient preoperative skin preparation: Helps to reduce bacteria that potentially can cause skin infection. ▶



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1 /Pouch

Made in China



D07310901 Rev2

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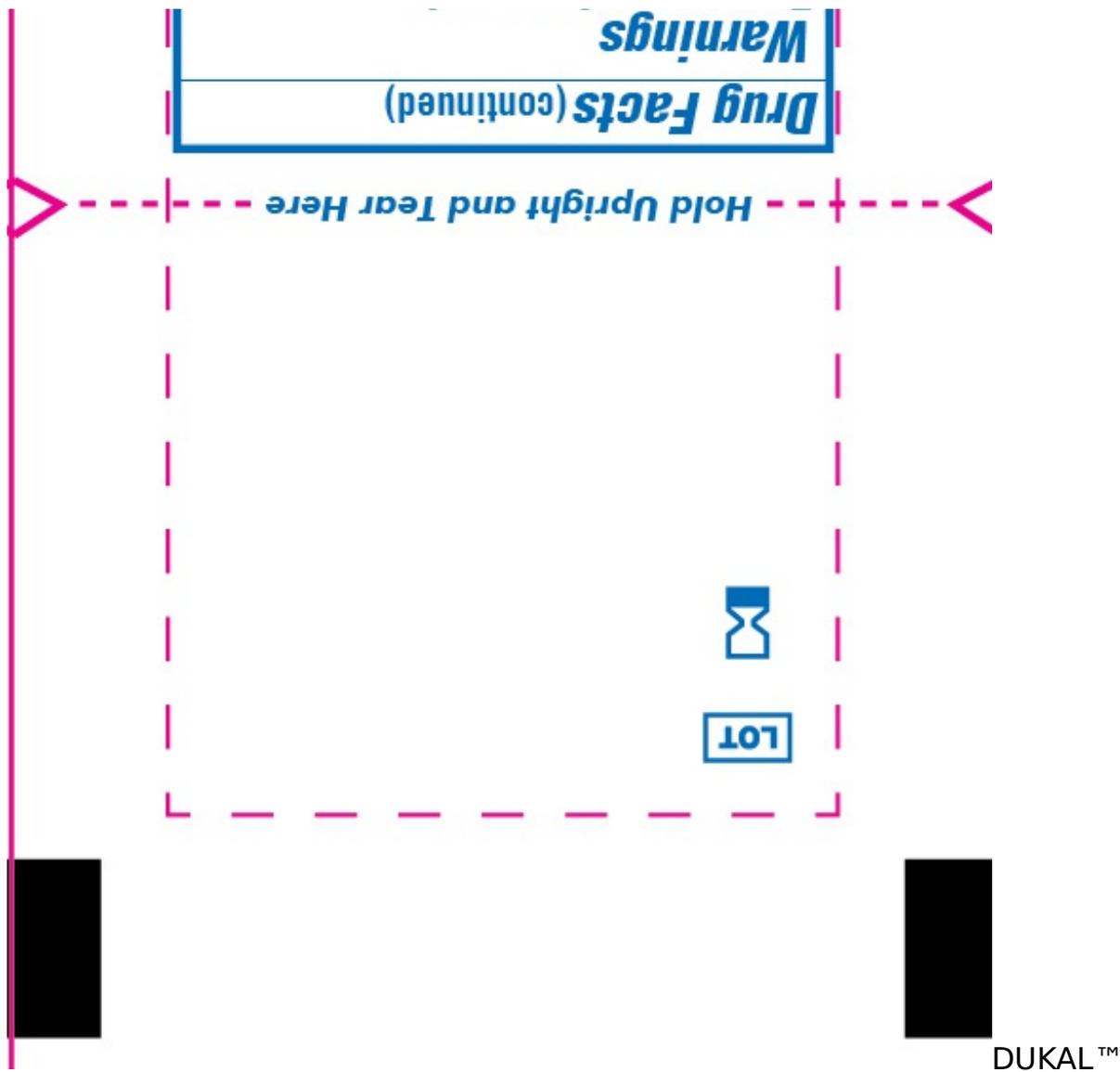
Inactive ingredients
alkyl glucoside, anhydrous citric acid,
sorbitol-10, glycerin, hydroxyethyl
cellulose, potassium iodine, sodium
hydroxide, water

Questions? 1-800-243-0741

Manufactured For:

DUKAL CORPORATION
Ronkonkoma, NY 11779

www.dukal.com



CORPORATION

NDC 65517-0036-1

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1/Pouch 2

Made in China D07310901 Rev2

POVIDONE-IODINE GEL SWABSTICK

povidone iodine gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:65517-0036
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 g

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
HYDROXYETHYL CELLULOSE (5000 CPS AT 1%) (UNII: X70SE62ZAR)	
POTASSIUM IODIDE (UNII: 1C4QK22F9J)	
GLYCERIN (UNII: PDC6A3C0OX)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
NONOXYNOL-10 (UNII: K7O76887AP)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:65517-0036-1	2.3 g in 1 POUCH; Type 0: Not a Combination Product	12/02/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003	12/02/2019	

Labeler - Dukal LLC (791014871)

Revised: 10/2023

Dukal LLC