# DUAL SWAB SCRUB SWABSTICK- povidone iodine swab Dynarex Corporation

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.

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#### **Dual Swab Povidone Iodine Scrub Swabstick**

#### **Active Ingredient**

Active ingredient Purpose Povidone iodine 7.5% Antiseptic

#### **Purpose**

For scrubbing procedure site, prior to antiseptic application

#### Indications and usage

For pre-operative patient skin preparation.

#### **Warnings**

#### **IFor External Use Only**

**Avoid contact with the eyes**. If contact occurs, flush with water.

#### Stop use

#### Stop use and ask a healthcare professional if,

- condition worsens or persists for more than 72 hours
- if pain, irritation, redness or swelling occurs

#### Ask a doctor

#### Ask a doctor before use if you have

- deep puncture wounds
- animal bites
- serious wounds

#### Do not use

#### Do not use:

- as a first aid antiseptic for more than 1 week
- over large areas of the body

#### Keep out of reach of children

#### Keep out of reach of children:

• if swallowed, get medical help or contact a Poison Control Center immediately.

#### Dosage and administration

#### **Directions**

#### For preoperative patient skin preparation:

• scrub procedure site as required. Discard after single use and proceed to packet 2.

#### As a first aid antiseptic:

- clean affected area
- apply to wound once or twice daily
- may be covered with a sterile bandage
- if bandaged, let dry first.

Tear at notch, remove applicator, use only once.

#### **Inactive ingredients**

Lauramide DEA, Water

#### Other information

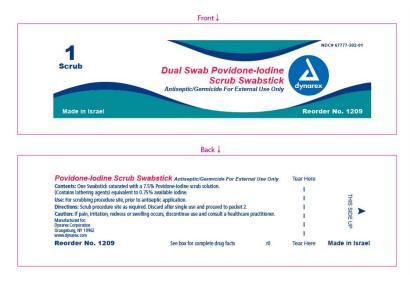
- Store at room temperature
- Avoid excessive heat

#### **Principal Display Panel**

Dynarex Dual Swab Povidone Iodine Scrub Swabstick

Item#: 1209

1209 Scrub.jpg



Active Ingredient	Purpose
	Antiseptio
Povidone Iodine Prep 10% w/w	Antiseptio
Uses	
Povidone-Iodine Scrub Swabstick • For scrubbing pro	
Povidone-Iodine Prep Swabstick • For antiseptic appli	cation at procedure site.
Warnings	
For External Use Only.	
Avoid contact with the eyes. If contact occurs	
Do not use • As a first aid antiseptic for more than	
Ask a doctor before use if you have • Deep	
Stop use and ask a healthcare practitioner • Irritation and redness develops	· if • Condition worsens or persists for more than 72 hours
Keep out of reach of children • If swallowed,	get medical help or contact a Poison Control Center immediately
Directions	
For preoperative patient skin preparation	
	te as required. Discard after single use and proceed to packet 2.
Povidone-Iodine Prep Swabstick • Apply to procedure immediate venipuncture, cover the area with a dry ste	site and allow to dry. Discard after single use. If not ready for rile gauze.
As a first aid antiseptic . Clean affected area .	
• May be covered with a sterile bandage • If bandage	d, let dry first
Tear at notch • Remove applicator • Use only once	
Other Information • Store at room temperature	Avoid excessive heat
Inactive Ingredients • Povidone Iodine 7.5% Sci	ub Solution – Lauramide DEA, water
<ul> <li>Povidone Iodine 10% Pre</li> </ul>	p Solution – Anhydrous citric acid, disodium hydrogen phosphate, wat

# DUAL SWAB SCRUB SWABSTICK povidone iodine swab Product Information Product Type HUMAN OTC DRUG Item Code (Source) Route of Administration TOPICAL

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
PO VIDO NE-IO DINE (UNII: 85H0 HZU99M) (IO DINE - UNII:9679TC07X4)	IO DINE	7.5 g in 100 g	

Inactive Ingredients		
Ingredient Name	Strength	
LAURIC DIETHANO LAMIDE (UNII: 12912VHG38)		
WATER (UNII: 059QF0KO0R)		

	Packaging				
ı	#	Item Code	Package Description	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
ı	1	1 NDC:67777-302-01 0.3 g in 1 POUCH; Type 0: Not a Combination Product			

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part333C	11/15/2015	

### Labeler - Dynarex Corporation (008124539)

## Registrant - Dynarex Corporation (008124539)

Establishment					
Name	Address	ID/FEI	Business Operations		
Sion Biotext Medical Ltd		532775194	manufacture(67777-302)		

Revised: 11/2015 Dynarex Corporation