POVIDONE-IODINE PREP PAD - povidone-iodine swab Yinjing Medical Technology (Shanghai) Co., Ltd.

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 PREP PAD

Drug Facts

Active Ingredients

Povidone-Iodine 10% w/w (1% titratable iodine)

Purpose

Antiseptic

Use

First aid antiseptic to help prevent infection in scrapes, minor cuts and burns Antiseptic to prepare skin prior to surgery. Uses antiseptic skin preparation

Stop Use Section

Stop use and ask a doctor if skin irritation, redness, swelling, or pain occurs.

Warnings Section

Warnings • Do not use if allergic to iodine • For external use only • Do not use in eyes • Avoid pooling beneath patient • In case of accidental ingestion, seek professional assistance or consult a poison control center immediately.

Keep Out of Reach of Children Section

Keep out of reach of children.

Directions Section

Directions apply locally as needed

Inactive Ingredients Section

Inactive ingredients Sodium hydroxide, water

Other Information Section

Other information For Hospital or Professional Use Only. Store at room temperature: 15 \mathbb{I} - 30 \mathbb{I}

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LOT EXP.

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NDC:44019-211-01

1 pad/pouch

POVIDONE-IODINE PREP PAD

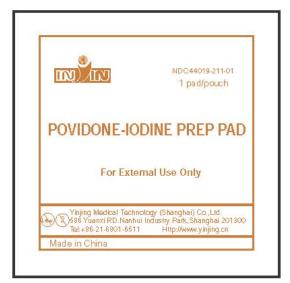
For External Use Only

Product Label

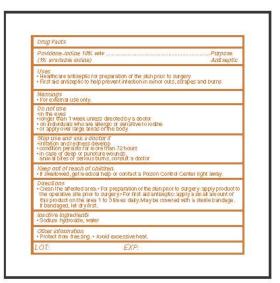


1525c

50mm



50_{mm}



POVIDONE-IODINE PREP PAD

povidone-iodine swab

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:44019-211
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:9679 TC07X4)	IODINE	10 g in 100 g	

Inactive Ingredients		
Ingredient Name	Strength	
SO DIUM HYDRO XIDE (UNII: 55X0 4QC32I)		
WATER (UNII: 059QF0KO0R)		

I	Packaging				
	# Item Code	Package Description	Marketing Start Date	Marketing End Date	
	NDC:44019-211-01	1 in 1 POUCH	08/03/2016		
	1	0.45 g in 1 PATCH; Type 0: Not a Combination Product			

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part333A	08/03/2016		

Labeler - Yinjing Medical Technology (Shanghai) Co., Ltd. (530501535)

Registrant - Yinjing Medical Technology (Shanghai) Co., Ltd. (530501535)

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