

EASYTOUCH ALCOHOL PREP PADS STERILE- isopropyl alcohol swab
MHC Medical Products, 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.

Easy•Touch® Alcohol Prep Pads, Sterile

Drug Facts

Active ingredient

Isopropyl Alcohol, 70% by volume

Antiseptic

Use

For preparation of the skin prior to injection

Warnings

For external use only. Flammable, keep away from fire or flame.

Do not use with electrocautery procedures, or in/near eyes.

Stop use if irritation or redness develops.

If irritating condition persists for more than 72 hours, consult a physician.

Keep out of reach of children. If swallowed, seek medical attention and/or contact a Poison Control Center immediately.

Directions

Prepare site by wiping vigorously

Inactive ingredient

Purified water

PRINCIPAL DISPLAY PANEL - 100 Medium 2-Ply Prep Pads Carton

Easy•Touch®

Alcohol Prep Pads

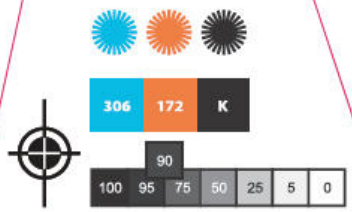
**Gamma-
Sterilized**

100 Medium 2-Ply Prep Pads

For External Antiseptic Use Only

Isopropyl Alcohol, 70% by Volume

Sterilized with Gamma Radiation



Easy•Touch®
Alcohol Prep Pads

Gamma-Sterilized

100 Medium 2-Ply Prep Pads

Easy•Touch®
Alcohol Prep Pads

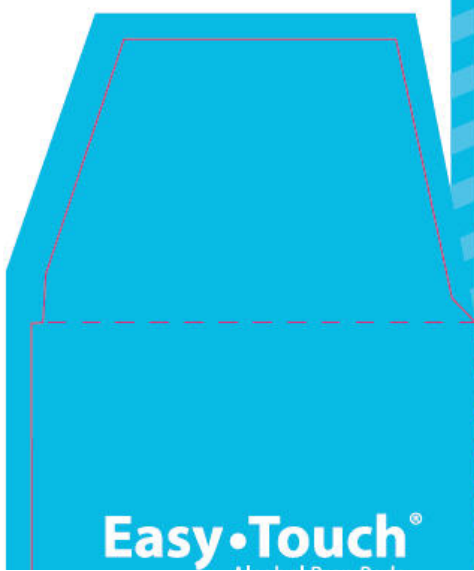
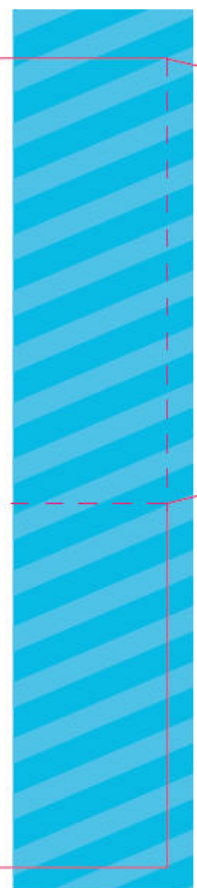
Gamma-Sterilized

100 Medium 2-Ply Prep Pads
For External Antiseptic Use Only
Isopropyl Alcohol, 70% by Volume
Sterilized with Gamma Radiation

2-Ply Prep Pads



ITEM #: 802711 LOT #:



Easy•Touch®

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For External Antiseptic Use Only
Isopropyl Alcohol, 70% by Volume
Sterilized with Gamma Radiation

Gamma-Sterilized

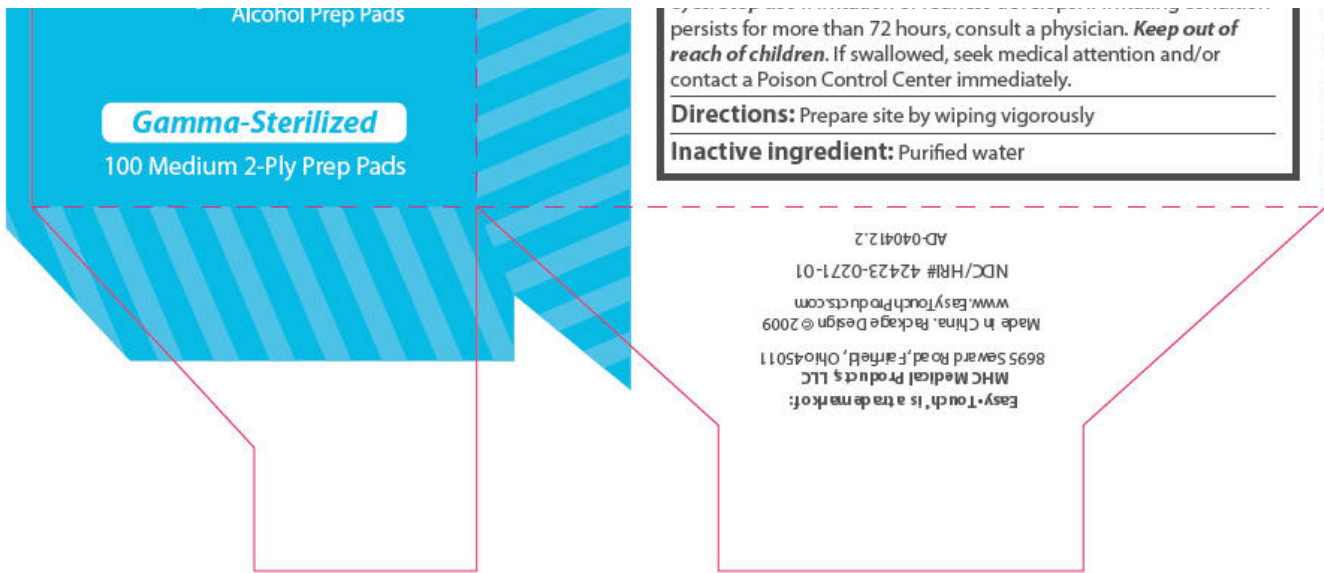
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EASYTOUCH ALCOHOL PREP PADS STERILE				
isopropyl alcohol swab				
Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:42423-271	
Route of Administration	TOPICAL			
Active Ingredient/Active Moiety				
Ingredient Name		Basis of Strength	Strength	
Isopropyl Alcohol (UNII: ND2M416302) (Isopropyl Alcohol - UNII:ND2M416302)		Isopropyl Alcohol	70 mL in 100 mL	
Inactive Ingredients				
Ingredient Name			Strength	
Water (UNII: 059QF0KO0R)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:42423-271-01	100 in 1 BOX	04/01/2014	
1		5 mL in 1 POUCH; Type 0: Not a Combination Product		
2	NDC:42423-271-02	200 in 1 BOX	04/01/2014	
2		5 mL 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 NOT FINAL	part333A	04/01/2014	

Labeler - MHC Medical Products, LLC (169895245)

Revised: 2/2023

MHC Medical Products, LLC