ALCOHOL PAD- alcohol prep pad swab Redicare 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.

Alcohol Prep Pads

Active Ingredient (in each towelette)Purpose

Isopropyl Alcohol 70%......Antiseptic

Use:

For preparation of skin prior to injection.

Warnings:

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

Do not use:

- With electrocautery procedures
- In the eyes

Stop use if

irritation and redness develop. If condition persists consult your health care practitioner.

Keep out of reach of children.

If swallowed, get medical help or contact a Poison COntrol Center right away.

Directions:

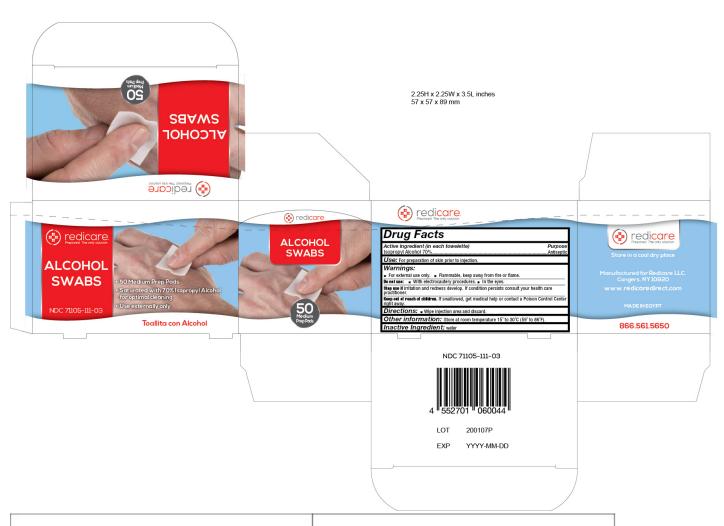
Wipe injection area and discard.

Other information:

Store at room temperature 15 ° to 30 °C (59 ° to 86 °F).

Inactive ingredient:

water





Drug Facts Active Ingredients Purpose Isopropyl Alcohol 70%.. .Antiseptic Use: For preparation of skin prior to injection. Warnings: . For external use only. Flammable, keep away from fire or flame. Do not use: With electrocautery procedures. In the eyes, If contact occurs flush eyes with water. Stop use if irritation and redness develop, if condition continues, consult your health care practitioner. Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away. Directions: . Wipe injection area and discard. Other Information: Store at room temperature 15° to 30°C (59" to 86°F). Inactive Ingredient: water LOT:

ALCOHOL PAD

alcohol prep pad swab

Product Information					
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:71105-111		
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 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
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WATER (UNII: 059QF0KO0R)

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71105-111-03	50 in 1 BOX	03/31/2017	
1	NDC:71105-111-02	1 in 1 PACKET		
1	NDC:71105-111-01	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	03/31/2017					

Labeler - Redicare LLC (800149346)

Revised: 12/2019 Redicare LLC