

ALCOHOL PREP- isopropyl alcohol swab
Acme United 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.

Acme Sterile Alcohol Prep Pads

Active Ingredient	Purpose
Isopropyl Alcohol 70% v/v	Antiseptic

Purpose

For preparation of the skin prior to injection.

Warnings:

- For external use only
- Flammable, keep away from flame or fire
- Not for use with electrocautinary devices or procedures
- Do not use in eyes
- Sterile unless package is damaged or open.

Indications and Usage:

Stop use and ask a doctor if:

- Irritation or redness develops
- condition persists for more than 72 hours
- Cleansing of an injection site

Keep out of reach of children.

In case of accidental ingestion, seek professional assistance or consult a poison control center immediately.

Directions:

Wipe injection site vigorously and discard

Other information:

- Store at room temperature: 15 deg C to 30 deg C 59 deg F to 86 deg F
- avoid excessive heat

Inactive Ingredient

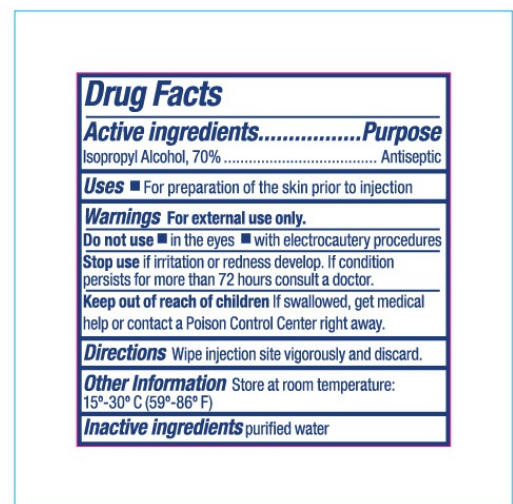
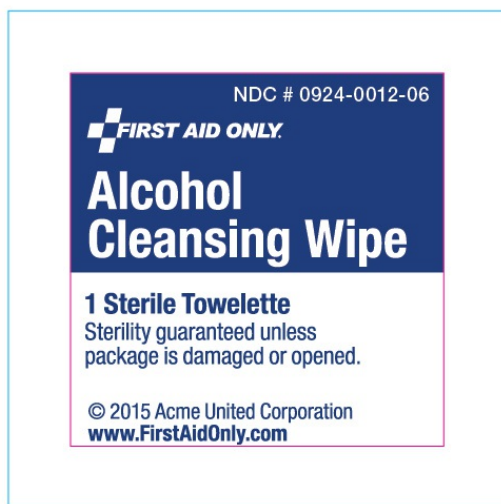
Inactive Ingredient

- Water

Principal Display Panel - Box



Principal Display Panel Pouch



ALCOHOL PREP

isopropyl alcohol swab

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0924-0012
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ISOPROPYL ALCOHOL (UNII: ND2M416302) (ISOPROPYL ALCOHOL - UNII:ND2M416302)	ISOPROPYL ALCOHOL	0.7 mL in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0924-0012-06	100 in 1 BOX	01/15/2016	
1		0.55 mL in 1 PACKET; 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	01/15/2016	

Labeler - Acme United Corporation (001180207)

Registrant - Dynarex Corporation (008124539)

Revised: 1/2016

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