

**FIRST AID ONLY ALCOHOL ANTISEPTIC- isopropyl alcohol liquid**

**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.*

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**First Aid Only Alcohol Antiseptic Pad**

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**Active ingredient**

Isopropyl Alcohol, 70% v/v

**Purpose**

Antiseptic Cleanser

**Use**

For preparation of the skin prior to an 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 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 site vigorously and discard.

**Other information**

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

**Inactive ingredient**

Purified Water



# Alcohol Antiseptic Pad

**1 Sterile Wipe**  
Sterility guaranteed unless package is damaged or opened.

Manufactured for:  
© 2020 Acme United Corporation  
55 Walls Dr., Fairfield, CT 06824 [www.FirstAidOnly.com](http://www.FirstAidOnly.com)  
Made in China 807440-revA

## Drug Facts

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Isopropyl Alcohol, 70% .....	Antiseptic Cleanser

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**Inactive ingredient** purified water

LOT

EXP

## FIRST AID ONLY ALCOHOL ANTISEPTIC

isopropyl alcohol liquid

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:0924-0017(NDC:59050-444)
<b>Route of Administration</b>	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:0924-0017-01	0.4 mL in 1 POUCH; Type 0: Not a Combination Product	06/15/2020	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date

OTC monograph not final	part333A	06/15/2020	
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**Labeler** - Acme United Corporation (001180207)

**Establishment**

Name	Address	ID/FEI	Business Operations
Acme United Corporation		045924339	relabel(0924-0017) , repack(0924-0017)

**Establishment**

Name	Address	ID/FEI	Business Operations
Acme United Corporation		080119599	relabel(0924-0017) , repack(0924-0017)

Revised: 6/2020

Acme United Corporation