

ALCOHOL PREP PAD- isopropyl alcohol swab
Tongzhou Deqi Medical Products Factory

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.

Drug Facts

Active Ingredient

Isopropyl Alcohol 70%

Purpose

Antiseptic

Use

For preparation of skin prior to injection prior to injection and to decrease germs in minor cuts and scrapes,

Warning

For External use only.

Stop use and ask a doctor if irritation develops.

Keep out of reach of children except with adult supervision.

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

Directions

Apply topically as needed to cleanse intended area.

Discard after single use.

Inactive Ingredients

Inactive Ingredients: Water

==== **STRICT**[®] ====

NDC: 71222-001-01

Alcohol

ALCOHOL Prep Pad

70% Isopropyl Alcohol

For Professional and Hospital Use

Tongzhou Deqi Medical Products Factory

No.1, Qibei Village, Tongzhou District, Nantong, Jiangsu, China

Drug Facts

Active ingredient

Isopropyl Alcohol 70%

Purpose

Antiseptic

Use for the preparation of skin prior to injection and to decrease germs in minor cuts and scrapes.

Non-sterile Solution
Applicator is sterile if package is intact

Warnings

For external use only.

Stop use and ask a doctor if irritation develops.

Keep out of reach of children except with adult supervision. If swallowed, get medical help or contact a Poison Control Center right away.

Directions

- Apply topically as needed to cleanse intended area.
- Discard after single use.

Inactive ingredient Purified water.

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ALCOHOL PREP PAD

isopropyl alcohol swab

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:71222-001
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

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	0.3 mL

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71222-001-01	1 in 1 PACKAGE; Type 0: Not a Combination Product	05/01/2016	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	05/20/2013	

Labeler - Tongzhou Deqi Medical Products Factory (544464252)

Establishment

Name	Address	ID/FEI	Business Operations
Tongzhou Deqi Medical Products Factory		544464252	manufacture(71222-001)

Revised: 2/2017

Tongzhou Deqi Medical Products Factory