

ALCOHOL PREP PAD- isopropyl alcohol liquid
Global Healthcare

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 Pad (6.5cm x 3cm)

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Active Ingredient

Isopropyl Alcohol, 70% v/v

Purpose

Antiseptic

Use

For preparation of skin prior to injection

Warnings

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

Do not use

*with electro cautery procedures * in the eyes.

Stop Use

if irritation or redness develop. If your condition persists for more than 72 hours, consult a doctor.

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 after single use.

Inactive ingredient

purified water

NDC No.: 42947-XXXXX

Alcohol Prep Pad

One Prep Pad (6.5cm x 3cm)

Saturated with 70% Isopropyl Alcohol

STERILE G Sterile if unopened or undamaged.

Antiseptic for preparation of the skin prior to an injection For External Use Only Discard After Use

2 Single use 1 piece/uniadd Item # 25P-003

LATEX - FREE

SIN - LATEX

ghc Global Healthcare

Manufactured in PRC for:

Global Healthcare - USA

Alpharetta, GA 30009

www.globalhealthcare.net

Alcohol Prep Pad (6.5cm x 3cm) 1 Pad (59294-000-00)

NDC No.: 42947-XXXXX

Alcohol Prep Pad

**One Prep Pad (6.5cm x 3cm)
Saturated with 70% Isopropyl Alcohol**



Sterile if unopened or undamaged.
Estéril si no está abierto o dañado.



Single use
Uso único

LATEX · FREE
SIN · LATEX



Antiseptic for preparation
of the skin prior to an injection
For External Use Only
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1 piece/unidad
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Drug Facts

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ALCOHOL PREP PAD

isopropyl alcohol liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:59294-000
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
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:59294-000-00	1 in 1 PACKET	07/11/2013	
1		0.4 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	07/11/2013	

Labeler - Global Healthcare (884718776)**Establishment**

Name	Address	ID/FEI	Business Operations
Wuxi Medical Instrument Factory		421292863	manufacture(59294-000)

Revised: 1/2020

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