

VIAMED ALCOHOL PREP PAD- alcohol swab
Rece International Corp

VIAMED Alcohol Prep Pad

Drug facts

Active Ingredient

Isopropyl Alcohol 70%

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 electrocautery 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 the reach of children

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

Directions

Wipe injectio site vigorously and discard after single use.

Inactive Ingredient

Purified water

Product labeling

ViaMed

Alcohol Prep Pad

For external use only

One pad Saturated with 70% Isopropyl Alcohol

Made in China

Rece International Corp

Miami Lakes, FL 33014

NDC No.:

VIA MED®

ISO9001

CE 0197

19710

ALCOHOL PREP PAD
TOALLITAS ALCOHOLADAS

FOR EXTERNAL USE ONLY
SOLO PARA USO EXTERNO

One Pad Saturated With 70% Isopropyl Alcohol
Una Toallita Impregnada en Alcohol Isopropílico al 70%

Made In China

RECE INTERNATIONAL CORP.

MIAMI LAKES, FL.33014

Drug Facts

Active Ingredient Purpose

Isopropyl Alcohol, 70% v/v.....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 electrocautery 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

LOT 150625

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VIAMED ALCOHOL PREP PAD

alcohol swab

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:70006-500
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	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:70006-500-01	100 in 1 PACKAGE	09/10/2015	
1		0.36 g 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 Drug	505G(a)(3)	09/10/2015	

Labeler - Rece International Corp (787828537)

Registrant - Wuxi Medical Instrument Factory (421292863)

Revised: 10/2023

Rece International Corp