ALCOHOL PREP PAD- isopropyl alcohol swab
NDC National Distribution & Contracting, Inc.

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.

Sterile

Alcohol Prep Pads

Drug Facts

Active Ingredients

Isopropyl Alcohol 70%

Purpose

Antiseptic Cleanser

Use

For Preparation of 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

PRINCIPAL DISPLAY PANEL - 100 Pouch Box Label

REF: **P902050**

Sterile

Alcohol
Prep Pads

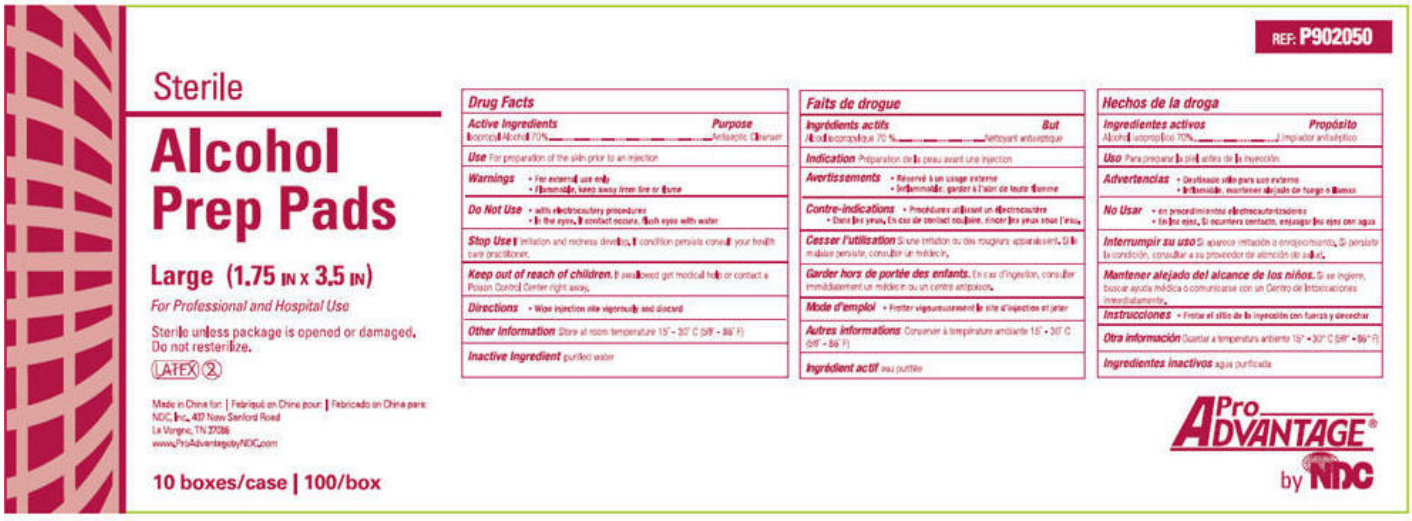
Large (1.75 IN × 3.5 IN)

For Professional and Hospital Use

Sterile unless package is opened or damaged.
Do not resterilize.

Pro
Advantage ®
by **NDC**

10 boxes/case | 100/box



ALCOHOL PREP PAD

isopropyl alcohol swab

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:43128-050
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:43128-050-01	100 in 1 BOX	12/15/2011	
1		1 mL in 1 POUCH; Type 0: Not a Combination Product		
2	NDC:43128-050-02	200 in 1 BOX	12/15/2011	
2		1 mL in 1 POUCH; 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	12/15/2011	

Labeler - NDC National Distribution & Contracting, Inc. (009831413)

Establishment

Name	Address	ID/FEI	Business Operations
Dukal		421317073	manufacture(43128-050)

Revised: 3/2023 NDC National Distribution & Contracting, Inc.