ISOPROPYL ALCOHOL ANTISEPTIC PAD- isopropyl alcohol swab Total Resources International 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.

Alcohol Prep Pads

Active Ingredients

Isopropyl Alcohol 70% v/v

Purpose

Antiseptic

Use

For the preparation of the skin prior to 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

Keep out of reach of children

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

Directions

Wipe Injection site vigorously and discard

Inactive Ingredient

purified water

Alcohol Prep Pad

1 Pc

One time use only



ISOPROPYL ALCOHOL ANTISEPTIC

Contents of unopened, undamaged package are sterile Made in USA * Manufactured for Total Resources Intl. Inc. Walnut, California 91789 * www.besmartgetprepared.com 00-ALC-90730 Rev.03 • NDC #55550-261-04





Active ingredient Isopropyl Alcohol 70% v/v Purpose Antiseptic

Use For preparation of the skin prior to 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 develops, If condition persists consult your healthcare practitioner

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control center immediately

Directions Wipe injection site vigorously and discard

Inactive ingredient purified water

ISOPROPYL ALCOHOL ANTISEPTIC PAD

isopropyl alcohol swab

Product Information

NDC:55550-261 **Product Type HUMAN OTC DRUG** Item Code (Source)

TOPICAL Route of Administration

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
9.04.0	

WATER (UNII: 059QF0KO0R)

Packaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
	NDC:55550-261- 04	1.4 mL in 1 POUCH; Type 0: Not a Combination Product	08/04/2022		

Marketing In	arketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part333E	08/04/2022		

Labeler - Total Resources International Inc. (790160535)

Revised: 8/2022 Total Resources International Inc.