MED-NAP STERILE ALCOHOL PREP PADS- isopropyl alcohol swab
Acme United Corporation

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.

MED NAP STERILE ALCOHOL PREP PADS

Drug Facts

Active ingredient

Isopropyl Alcohol 70% v/v

Purpose

Antiseptic Cleanser

Use For preparation of the skin prior to an injection

Do Not Use **● with electrocautery procedures ● In the eyes. If contact occurs, flush**

eyes with water

Slop Use If irritation or redness develop. If condition persists consult your health care practitioner.

Keep out of reach of children**.** If swallowed get medical help or consult a Poison

Control Center right away.

Directions **● Wipe injection site vigorously and discard.**

Other Information Sterile unless pouch is opened or damaged. Store at room temperature 15 º - 30 º C (59 º - 86 º F)

Inactive Ingredient purified water

Warnings**● For external use only ● Flammable, keep away from fire or flame**

Questions 800-835-2263

Prep Pad Box

MED-NAP STERILE ALCOHOL PREP PADS

isopropyl alcohol swab

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0924-0500
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**
	NDC:0924			

1	NDC:0924-0500-01	200 in 1 BOX	07/12/2021	
1		1 mL in 1 POUCH; Type 4: Device Coated/Impregnated/Otherwise Combined with Drug		
2	NDC:0924-0500-00	1 mL in 1 POUCH; Type 4: Device Coated/Impregnated/Otherwise Combined with Drug	07/12/2021	
3	NDC:0924-0500-02	50 in 1 BOX	07/12/2022	
3		1 mL in 1 POUCH; Type 4: Device Coated/Impregnated/Otherwise Combined with Drug		
4	NDC:0924-0500-04	100 in 1 BOX	07/12/2022	
4		1 mL in 1 POUCH; Type 4: Device Coated/Impregnated/Otherwise Combined with Drug		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	07/12/2021	

Labeler - Acme United Corporation (001180207)

Establishment

Name	Address	ID/FEI	Business Operations
Acme United Corporation		045924339	label(0924-0500) , pack(0924-0500)

Establishment

Name	Address	ID/FEI	Business Operations
Acme United Corporation		080119599	label(0924-0500) , pack(0924-0500)

Establishment

Name	Address	ID/FEI	Business Operations
Acme United Corporation		117825595	manufacture(0924-0500)

Revised: 7/2022 Acme United Corporation