# CERTUS ALCOHOL PREP- is opropyl alcohol swab Certus Medical, 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.

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## Certus Alcohol Prep Pad 210000 Drug Facts and Label

## **Drug Facts Box OTC-Active Ingredient Section**

Isopropyl Alcohol 70% v/v

## **Drug Facts Box OTC-Purpose Section**

Antiseptic

### **Drug Facts Box OTC-Indications and Usage Section**

For preparation of the skin prior to an injection

## **Drug Facts Box OTC-Warnings Section**

for external use only

flammable, keep away from fire or flame

## **Drug Facts Box OTC-Do Not Use Section**

with electrocautery procedures

in the eyes

if contact occurs, flush eyes with water

## **Drug Facts Box OTC-Stop Use Section**

if irritation and redness develop

if condition persists consult your health care practitioner

#### Drug Facts Box OTC-Keep Out Of Reach Of Children Section

if swallowed get medical help or call a Poison Control Center right away

## Drug Facts Box OTC-Dosage & Administration Section

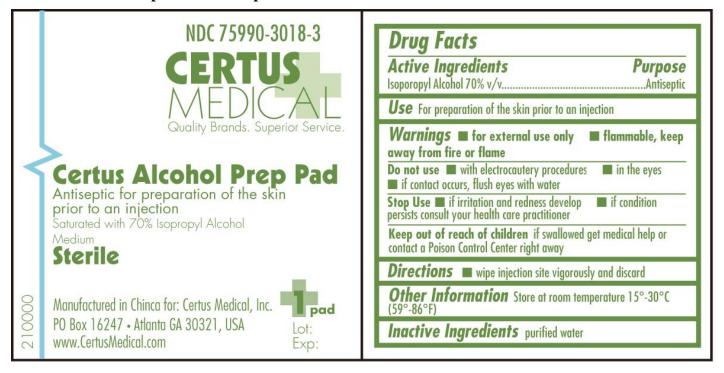
wipe injection site vigorously and discard

### **Drug Facts Box OTC-General Precautions Section**

Store at room temperature 15 degrees - 30 degrees C (59 degrees - 86 degrees F)

#### **Drug Facts Box OTC-Inactive Ingredient Section**

## Certus Alcohol Prep Pad 210000 pouch



## Certus Alcohol Prep Pad

#### CERTUS ALCOHOL PREP

CERT US ALCOHOL PREP								
isopropyl alcohol swab								
Product Information								
Product Type	HUMAN OTC DRUG	Item Code (Source)		NDC:75990-3018				
Route of Administration	TOPICAL							
Active Ingredient/Active Moiety								
Ing	gredient Name		Basis of S	Strength	Strength			
ISOPROPYL ALCOHOL (UNII: ND2M UNII: ND2M416302)	416302) (ISOPROPYL ALCOHO	L -	ISOPROPYL ALCOHOL		0.7 mL in 1 mL			

ı	Inactive Ingredients	
	Ingredient Name	Strength
	WATER (UNII: 059QF0KO0R)	

Packaging							
# Item Code	Package Description	<b>Marketing Start Date</b>	<b>Marketing End Date</b>				
1 NDC:75990-3018-3	0.4 mL in 1 POUCH; Type 0: Not a Combination Product	05/01/2011					

Marketing Information							
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date				
OTC monograph not final	part333A	05/01/2011					

Labeler - Certus Medical, Inc. (966433653)

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