

**SURE COMFORT ALCOHOL PREP PADS- isopropyl alcohol liquid
IDO PHARM**

isopropyl alcohol 70%

Water

apply topically as needed to cleanse

Keep out of reach of children

wipe injection site vigorously and discard after single use

Flammable, keep away from fire and flames

For external use only

For external use only

<p>SureComfort Alcohol Prep Pad 26-1260</p>	 <p>allison medical inspired innovation</p>	<ul style="list-style-type: none"> • www.allisonmedical.com • 1-800-886-1618 • 8091 Shaffer Parkway, Littleton, CO 80127
<p>Project: Alcohol Prep Pad Description: APP Single Pack Item#: 26-1260 NRC#: 86227-0221-58 Die#: 1.9375 x 1.9375 inches Vendor: Korea Date Released: Nov., 2024 PrMgr/Dir/Dsr: AM:CM/CC:SR Part#: 13387-101-01 Units Per Box: 1 count</p>	<p>PMS Spot Color</p> 	<p>4 Color Match </p>  <p>PMS 185 KEY LINE does NOT print Match color M 91:Y 76</p>

LOT		
Drug Facts		
Active Ingredient	Purpose	
Isopropyl Alcohol, 70% v/v	Antiseptic Cleaner	
Use	Apply topically as needed to cleanse	
Warnings: For external use only.		
Do not use - with electrocautery procedures - in the eyes.		
Stop use. Ask a doctor if irritation or redness develop.		
Flammable , keep away from fire or flame.		
Keep out of reach of children: If swallowed, seek medical help immediately or contact Poison Control.		
Directions: Wipe injection site vigorously and discard after single use.		
Inactive ingredient:	Purified water	
		13287-101-01

REF 26-1260	Sterile
SureComfort <i>Delivering Quality and Comfort</i>	
Alcohol Prep Pads	
Isopropyl Alcohol, 70% v/v	
Antiseptic for preparation of the skin prior to an injection	
Contains One Pad	
For External Use Only Discard After Use	

SURE COMFORT ALCOHOL PREP PADS

isopropyl alcohol liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:77039-032
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ISOPROPYL ALCOHOL (UNII: ND2M416302) (ISOPROPYL ALCOHOL - UNII:ND2M416302)	ISOPROPYL ALCOHOL	70 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:77039-032-01	0.2 mL in 1 POUCH; Type 0: Not a Combination Product	11/27/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003	04/24/2024	

Labeler - IDO PHARM (694853523)

Revised: 11/2024

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