

**AMERICAN RED CROSS ALCOHOL PREP PAD- isopropyl alcohol swab
YIWU HAODING MEDICAL CO.,LTD**

69139-011 Alcohol prep pad

DRUG FACTS

Active Ingredient

Isopropyl Alcohol 70%

Purpose

Antiseptic

Uses:

For preparation of the skin prior to an injection

Use

Apply alcohol as needed to clean intended area. Discard after single use.

Warning

For external use only.

Flammable,keep away from heat, spark, electrical, fire, or flame.

Do not use:with electrocautery procedures or in eyes.

Stop useif redness or irritation develops. If condition persists for more than 72 hours, consult a physician.

Keep out of reach of children.

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

Inactive Ingredients:

water



Drug Facts	
Active ingredient	Purpose Antiseptic
Isopropyl Alcohol, 70%	
Inactive Ingredient ■ purified Water.	
Use ■ for preparation of the skin prior to an injection.	
Directions ■ Apply alcohol as needed to clean intended area. Discard after single use.	
Warnings	
For external use only.	
Flammable, keep away from heat, spark, electrical, fire or flame.	
Do not use ■ with electrocautery procedures or in eyes.	
Stop use ■ If redness or irritation develops. If condition continues for more than 72 hours, consult a physician.	
Keep out of reach of children ■ If ingested, do not induce vomiting. Get medical help or contact a Poison Control Center immediately.	
Storage ■ Keep at room temperature.	Expiry ■ Two years
Importer:MY IMPORTS USA LLC 60 Brunswick Avenue Edison, NEW JERSEY, 08817, UNITED STATES	
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AMERICAN RED CROSS ALCOHOL PREP PAD

isopropyl alcohol swab

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69139-011
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:69139-011-01	1 in 1 PACKAGE	04/30/2025	
1		0.2 mL in 1 APPLICATOR; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M004	04/30/2025	

Labeler - YIWU HAODING MEDICAL CO.,LTD (421362384)

Establishment

Name	Address	ID/FEI	Business Operations
YIWU HAODING MEDICAL CO.,LTD		421362384	manufacture(69139-011)

Revised: 4/2025

YIWU HAODING MEDICAL CO.,LTD