

ETHYL ALCOHOL- alcohol gel
ASP Global LLC

ETHYL ALCOHOL GEL

DRUG FACTS

Each 100ml Contains

Active ingredient

Ethyl Alcohol I.P. :

70.0% v/v Gel base

USES

Hand Sanitization to decrease bacteria on the skin.

For use when soap and water are not available.

WARNINGS

Flammable. Keep away from fire or flame.

For external use only. Recommended for repeated use.

WHEN USING THE PRODUCT

Do not use in or near the eyes.

In case of contact, rinse eyes thoroughly with water.

STOP USE AND ASK A DOCTOR IF

Irritation or rash appears, **Keep out of reach of children.** If swallowed, get medical help or call a poison control center.

DIRECTIONS

Not recommended for infants. Wet hands thoroughly with product & allow to self dry. Children under 6 years of age should be supervised when using this product.

OTHER INFORMATION

Avoid freezing, Do not store above 40°C (104°F).

INGREDIENTS

Ultrez - 20. Glycerin, Propylene Glycol, Ethyl Alcohol, Aloe Vera Juice, Fragrance &

Purified Water.

DISTRIBUTED BY:

ASP GLOBAL, LLC
7800 THIRD FLAG PARKWAY,
AUSTELL, GA 30168, USA

PRINCIPAL DISPLAY PANEL - 53 mL Bottle Label

ASP

GLOBAL

INSTANT

Hand

Antiseptic

70.0% Ethyl Alcohol

1.9 FL OZ / 53 mL



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**MADE IN INDIA
FOR EXPORT ONLY**

NDC NO. 59448-011-01
REF #: 330137 REV 00

MFG. LIC: MNB/07/590 | B.No, Mfg. Date, Exp.Date, see bottle

Neutral Code # HP/Drugs/07/140

ETHYL ALCOHOL

alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:59448-011
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	70 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
Water (UNII: 059QF0KO0R)	
Aloe Vera Leaf (UNII: ZY81Z83H0X)	
Glycerin (UNII: PDC6A3C0OX)	
Propylene Glycol (UNII: 6DC9Q167V3)	
ACRYLATES/C10-30 ALKYL ACRYLATE CROSSPOLYMER (60000 MPa.S AT 0.5%) (UNII: YY2H MJ9NZ F)	
FRAGRANCE 13576 (UNII: 5EM498GW35)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:59448-011-01	53 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/01/2025	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph drug	M003	03/02/2025	

Labeler - ASP Global LLC (080361159)

Revised: 4/2025

ASP Global LLC