HAND SANITIZER- alcohol gel
Be Moxe, LLC

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.

HAND SANITIZER

Drug Facts

Active ingredient[s]

Ethyl Alcohol 70% v/v

Purpose

Antiseptic

Use[s]

Hand sanitizer to help reduce bacteria that potentially can cause disease. For use when soap and water are not available.

Warnings

For external use only. Flammable. Keep away from heat or flame.

Do not use

- in children less than 2 years of age
- on open skin wounds

When using this product keep out of eyes, ears, and mouth. In case of contact with eyes, rinse eyes thoroughly with water.

Stop use and ask a doctor if irritation or rash occurs. These may be signs of a serious condition.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions

- Place enough product on hands to cover all surfaces. Rub hands together until dry.
- Supervise children under 6 years of age when using this product to avoid swallowing.

Other information

- Store between 59-86°F (15-30°C)

- Avoid freezing and excessive heat above 104°F (40°C)

Inactive ingredients

Water (Aqua), Glycerin, Isopropyl Myristate, Carbomer, Triethanolamine.

Distributed by:
Be MOXĒ LLC 4700 140th Ave N., Ste 112,
Clearwater, FL 33762

PRINCIPAL DISPLAY PANEL - 237 ml Bottle Label

MOXĒ

HAND
SANITIZER

70% Alcohol

MADE IN
USA

8FL OZ | 237ml

HAND SANITIZER

alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:75435-727
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)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	

TROLAMINE (UNII: 9O3K93S3TK)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:75435-727-08	237 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/01/2020	
2	NDC:75435-727-10	295 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/01/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH NOT FINAL	part333A	06/01/2020	

Labeler - Be Moxe, LLC (114624756)

Revised: 4/2022

Be Moxe, LLC