MOXE CITRUS 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

Ethyl Alcohol 70% by volume

Purpose

Antiseptic

Uses

Use to help reduce bacteria on the skin.

Warnings

- **For external use only.**
- Flammable.
- Keep away from fire or flame.

Do Not Use Near Eyes.

In case of contact, rinse eyes thoroughly with water.

Stop Use And Ask A Doctor

If skin irritation develops

Keep Out Of Reach Of Children.

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

Directions

- Place sanitizer on palm, rub hands together.
- Supervise children under 6.

Other Information

Do not store over 110° F

Questions?

Call 1-800-296-3160

Inactive Ingredients

Water, Carbomer, Glycerin, Aloe Leaf Juice, Tocopheryl Acetate, Fragrance

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

PRINCIPAL DISPLAY PANEL - 118 ml Bottle Label

MOXĒ

CITRUS

HAND
SANITIZER

70% Alcohol

MADE IN
USA

4FL OZ | 118ml

MOXE CITRUS HAND SANITIZER

alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:75435-813
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)	
Carbomer Homopolymer, Unspecified Type (UNII: 0A5MM307FC)	
Glycerin (UNII: PDC6A3C0OX)	
Aloe (UNII: V5VD430YW9)	
.Alpha.-Tocopherol Acetate (UNII: 9E8X80D2L0)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:75435-813-04	118 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/17/2020	
2	NDC:75435-813-08	237 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/17/2020	
3	NDC:75435-813-12	355 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/17/2020	
4	NDC:75435-813-32	946 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/17/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH NOT FINAL	part333A	06/17/2020	

Labeler - Be Moxe, LLC (114624756)

Establishment

Name	Address	ID/FEI	Business Operations
JW Nutritional, LLC		017642837	MANUFACTURE(75435-813)

Revised: 2/2022 Be Moxe, LLC