

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.

MOXĒ CITRUS HAND SANITIZER

Drug Facts

Active Ingredient [s]

Alcohol w/w 70%

Purpose

Antiseptic

Use[s]

Use to help reduce bacteria on the skin.

Warnings

- **For external use only.**
- Flammable.
- Keep away from heat 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 years of age.

Other Information

Do not store over 110° F

Questions?

Call 1-800-296-3160

Inactive Ingredients

Water (Aqua), Glycerin, Propylene Glycol, Carbomer, Triethanolamine, Fragrance

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

PRINCIPAL DISPLAY PANEL - 89 ml Bottle Label

MOXĒ
 CITRUS
 HAND
 SANITIZER
 70% Alcohol
 MADE IN
 USA
 3FL OZ | 89ml



| MOXE CITRUS HAND SANITIZER | | | |
|--|----------------|--------------------|-----------------|
| alcohol gel | | | |
| Product Information | | | |
| Product Type | HUMAN OTC DRUG | Item Code (Source) | NDC:75435-003 |
| 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) | | | |

Glycerin (UNII: PDC6A3C0OX)

Propylene Glycol (UNII: 6DC9Q167V3)

Carbomer Homopolymer, Unspecified Type (UNII: 0A5MM307FC)

Trolamine (UNII: 9O3K93S3TK)

Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|------------------|--|----------------------|--------------------|
| 1 | NDC:75435-003-03 | 89 mL in 1 BOTTLE; Type 0: Not a Combination Product | 08/12/2020 | |

Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|-------------------------|--|----------------------|--------------------|
| OTC MONOGRAPH NOT FINAL | part333A | 08/12/2020 | |

Labeler - Be Moxe, LLC (114624756)

Revised: 8/2020

Be Moxe, LLC