MOXIE HAND SANITIZING GEL- alcohol gel
Skaffles Group

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

Moxie Hand Sanitizer Gel

Drug Facts

Active Ingredient

Ethyl Alcohol 71% V/V

Purpose

Antiseptic

Use

- for hand sanitizing to decrease bacteria on the skin
- recommended for repeated use

Warnings

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

When using this productdo not use in or near the eyes. In case of contact, rinse eyes thoroughly with water

Discontinue Use if irritation and redness develop. If condition persists for more than 72 hours consult a doctor

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

Directions

- Pump a small amount in your palm and briskly rub hands together until dry.
- Children under 6 years of age should be supervised when in use.

Other Information

Store between 15-30C (59-89F)

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

Inactive ingredients

Water, Glycerin, Carbomer, Triethanolamine, Aloe Barbadensis Leaf Juice, Tocopheryl Acetate (Vitamin E).

Ditributed by: Lowe's Home Centers LLC

Mooresville NC 28117

Principal Display Panel

Moxie

HAND SANITIZER GEL

Kills 99.9% of most common germs

Moisturizes with Aloe & Vitamin E

Light & Fresh Scent

16.9 FL OZ (500ml)

MADE IN CHINA

Item

Model

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MOXIE HAND SANITIZING GEL
alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:77720-033

Route of Administration	TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	71 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED) (UNII: 59TL3WG5CO)	
TROLAMINE (UNII: 9O3K93S3TK)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
ALPHA-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:77720-033-01	500 mL in 1 PACKAGE; Type 0: Not a Combination Product	05/15/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		05/15/2024	

Labeler - Skaffles Group (831115642)

Revised: 3/2025 Skaffles Group