PURPLE MOUNTAIN CLEAN HAND SANITIZER- isopropyl alcohol liquid Edison Nation, Inc.

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.

PURPLE MOUNTAIN CLEAN HAND SANITIZER

DRUG FACTS

Active Ingredients:

Isopropyl Alcohol 75% v/v

Purpose:

antimicrobial

Uses:

Hand sanitizer to help reduce bacteria that potentially can cause disease. For use when soap and water are unavailable.

Warnings:

For external use only. Flammable. Keep away from heat and flame, avoid contact with eyes. In case of contact with eyes, flush thoroughly with water. Avoid contact with broken skin. Stop use and ask doctor if irritation or redness develops, or if condition persists for more than 72 hours

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. For children under 6, use with adult supervision. Not recommended for infants.

Other info:

Store between 15-30C (59-86F).

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

Inactive Ingredients:

Glycerin, hydrogen peroxide, water USP, Fragrance

Refreshing Liquid

Made in the USA

Kills more than 99.9% of germs

75% alcohol

100% SATISFACTION

GUARANTEED

Distributed by:

GLOBAL CLEAN SOLUTIONS

WARNING

Packaging

PURPLE MOUNTAIN CLEAN HAND SANITIZER

isopropyl alcohol liquid

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:80120-102

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name Basis of Strength Strength

ISOPROPYL ALCOHOL (UNII: ND2M416302) (ISOPROPYL ALCOHOL - UNII: ND2M416302) ISOPROPYL ALCOHOL 75 L in 100 L

Inactive Ingredients

Ingredient Name Strength

GLYCERIN (UNII: PDC6A3C0OX)
HYDROGEN PEROXIDE (UNII: BBX060AN9V)

WATER (UNII: 059QF0KO0R)

Packaging

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	# Item Code	Package Description	Marketing Start Date	Marketing End Date		
	NDC:80120-102-00	3.785 L in 1 BOTTLE; Type 0: Not a Combination Product	08/17/2020			

Marketing Information

Harketing information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part333A	08/17/2020		

Labeler - Edison Nation, Inc. (081181069)

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