NINETEEN CERTIFIED HAND SANITIZER- alcohol liquid Lopez-Scripa Industries 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.

Nineteen Certified Hand Sanitizer

ETHYL ALCOHOL 80%

PURPOSE

Antiseptic

INACTIVE INGREDIENTS

17-18% Sterilized Water, 1.5% Glycerin 0.2% Hydrogen Peroxide 34%, Less than 0.01% Bittering Agent

USES

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

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. Do not use on children less than 2 months of age. Do not use on open wounds.

WARNINGS

FOR EXTERNAL USE ONLY - HANDS

FLAMMABLE - KEEP AWAY FROM HEAT AND FLAME

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 THE REACH OF CHILDREN

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

OTHER INFORMATION

Store between 59-86°F (15-30°C). Avoid freezing and excessive heat above 104°F (40°C). May discolor some fabrics. Harmful to wood finishes and plastics.

Package Labeling:



NINETEEN CERTIFIED HAND SANITIZER

alcohol liquid

right away.

ONLY - HANDS

Product Information

HUMAN OTC DRUG NDC:79526-000 Product Type Item Code (Source)

TOPICAL **Route of Administration**

Active Ingredient/Active Moiety

Ingredient Name Basis of Strength Strength ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M) ALCOHOL 0.8 mL in 1 mL

Inactive Ingredients

Ingredient Name Strength WATER (UNII: 059QF0KO0R) GLYCERIN (UNII: PDC6A3C0OX) HYDROGEN PEROXIDE (UNII: BBX060AN9V)

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Ш	Packaging						
	# Item Code	Package Description	Marketing Start Date	Marketing End Date			
П	NDC-70526 000	270 F. 41 m. in 1 POTTI E. Tyme O. Not a Combination					

1 01 NDC:/9520-000-	5/05.41 III. III. 1 BOTTLE; Type 0: Not a Combination Product	08/08/2020			
Marketing Information					
Marketing Catego	ry Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
OTC monograph not fi	nal part333E	08/08/2020			

Labeler - Lopez-Scripa Industries LLC (036115985)

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