HAND SANITIZER- alcohol liquid
Ver-tech Laboratories, 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.

Active Ingredient(s)

Alcohol 70% v/v

Purpose

Antiseptic

Use

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

Warnings

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

Do not use

on children less than 2 months of age

on open skin wounds

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 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. Supervise children under 6 years of age when using this product to avoid swallowing.

Other information

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

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

Inactive Ingredients

glycerin, hydrogen peroxide, purified water, fragrance

3.78 L NDC: 74807-970-01

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HAND SANITIZER

alcohol liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:74807-970
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**
CITRAL (UNII: T7EU0O9VPP)	
LEMON OIL (UNII: I9GRO824LL)	
HYDROGEN PEROXIDE (UNII: BBX060AN9V)	
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
LIME OIL (UNII: UZH29XGA8G)	
DIPROPYLENE GLYCOL (UNII: E107L85C40)	
METHACRYLIC ACID - ETHYL ACRYLATE COPOLYMER (4500 MPA.S) (UNII: T967IEU43C)	
MONOETHANOLAMINE (UNII: 5KV86114PT)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:74807-970-01	3780 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/04/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	05/04/2020	

Labeler - Ver-tech Laboratories, Inc. (166453097)

Establishment

Name	Address	ID/FEI	Business Operations
Ver-tech Laboratories, Inc.		166453097	manufacture(74807-970)

Revised: 5/2020 Ver-tech Laboratories, Inc.