MEDICAP HAND SANITIZER- alcohol gel
Zurich Pharma 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.

MEDICAP HAND SANITIZER

Drug Facts

Active ingredient[s]

Alcohol 66.05 % v/v

Purpose

Antiseptic

Use[s]

• 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

• in 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 15-30°C (59-86°F) • Avoid freezing and excessive heat above 40°C (104°F)

Inactive ingredients: Glycerine, Tetrahydroxypropyl Ethylenediamine, Hypromellose, Sodium Chloride, Acid Blue 25, Peppermint Essential Oil, Distilled Water, Citral, Dipropylene Glycol, Linalool, Citronellol, Coumarin, Isopropyl Alcohol.

Alcohol Antiseptic 66.05%

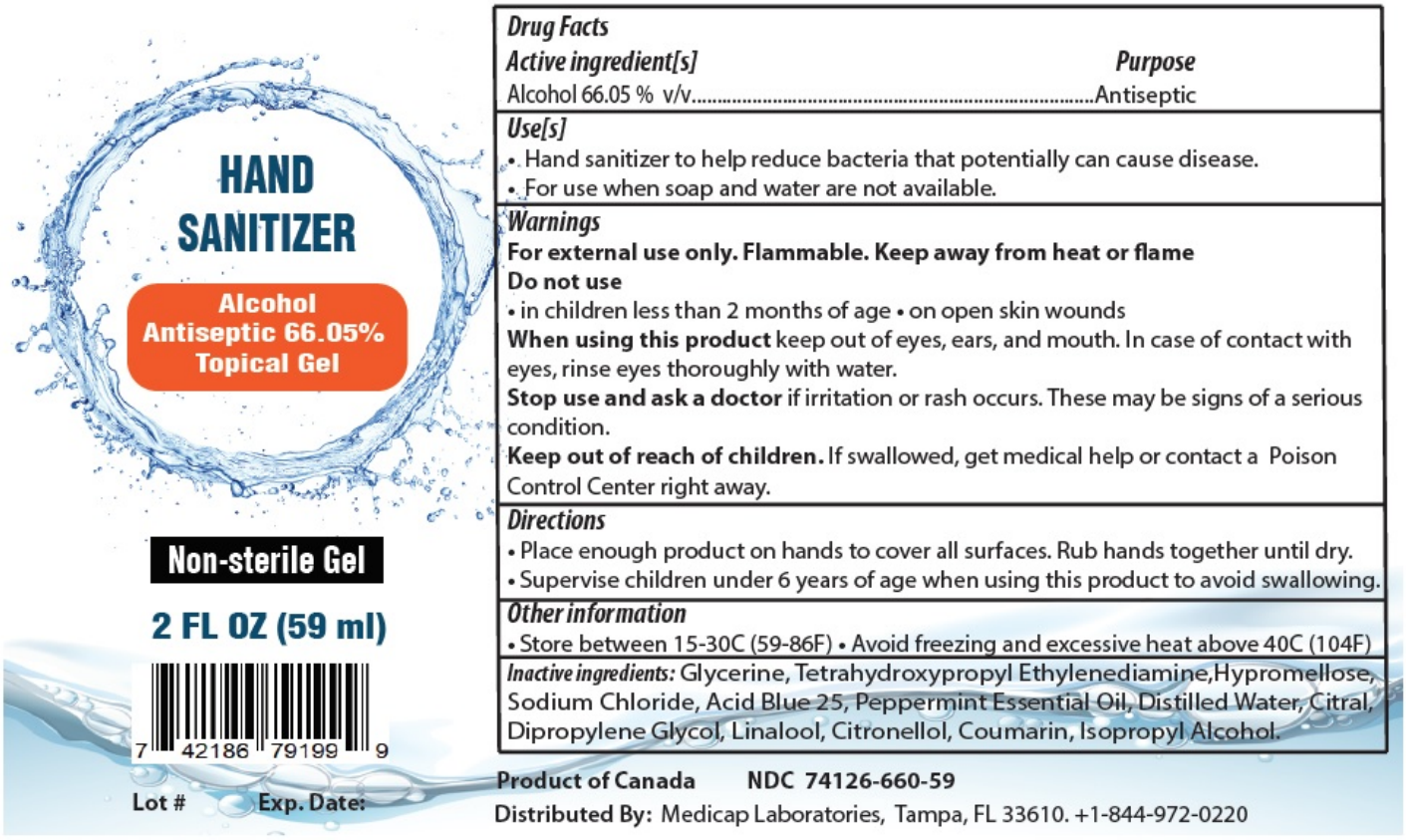
Topical Gel

Non-sterile Gel

Product of Canada

Distributed By: Medicap Laboratories, Tampa, FL 33610. +1-844-972-0220

Packaging



MEDICAP HAND SANITIZER

alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:74126-660
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	38.97 mL in 59 mL

Inactive Ingredients	
Ingredient Name	**Strength**

GLYCERIN (UNII: PDC6A3C0OX)	
EDETOL (UNII: Q4R969U9FR)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
ACID BLUE 25 (UNII: S2E15W6FSN)	
PEPPERMINT OIL (UNII: AV092KU4JH)	
WATER (UNII: 059QF0KO0R)	
CITRAL (UNII: T7EU0O9VPP)	
DIPROPYLENE GLYCOL (UNII: E107L85C40)	
LINALOOL, (+/-)- (UNII: D81QY6I88E)	
.BETA.-CITRONELLOL, (R)- (UNII: P01OUT964K)	
COUMARIN (UNII: A4VZ22K1WT)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:74126-660-59	59 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 - Zurich Pharma Inc (248184926)

Establishment

Name	Address	ID/FEI	Business Operations
Zurich Pharma Inc		248184926	manufacture(74126-660)

Revised: 5/2020 Zurich Pharma Inc