

**SANKIND HAND SANITIZER 75 % V/V ISOPROPYL ALCOHOL- isopropyl alcohol gel
UNIVET HOLDINGS LIMITED**

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.

Sankind Hand Sanitizer 75 % v/v Isopropyl Alcohol

Active Ingredient(s)

Isopropyl Alcohol 75% v/v. Purpose: Antiseptic

Purpose

Antiseptic, Hand Sanitizer

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

- 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. 1-800-222-1222

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.

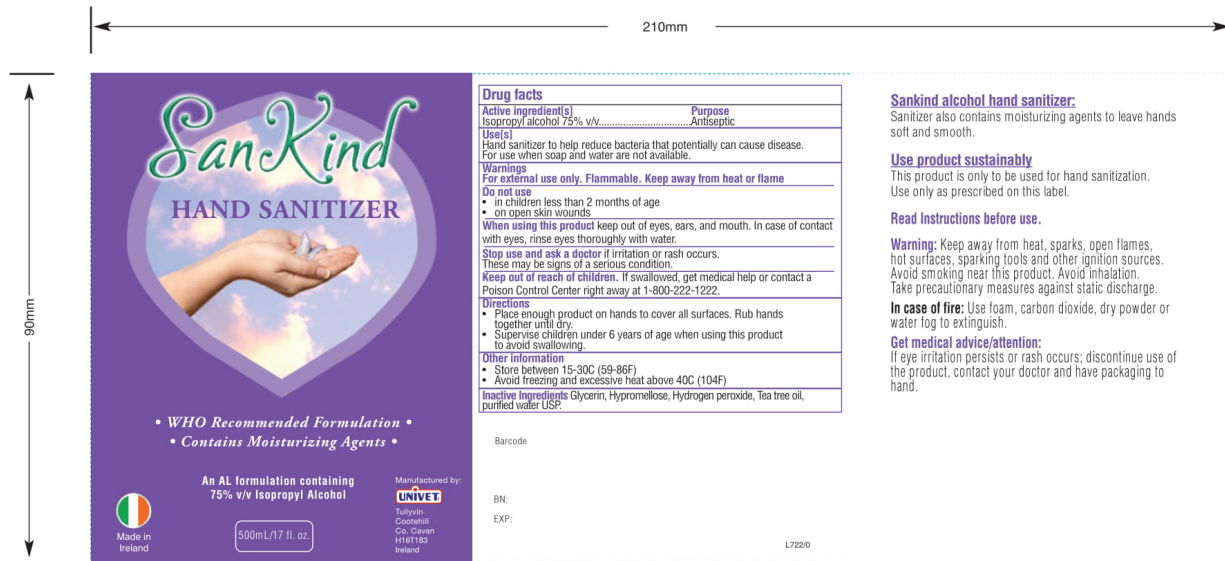
Other information

- Store between 15-30C (59-86F)
- Avoid freezing and excessive heat above 40C (104F)

Inactive ingredients

glycerin, Hypromellose, hydrogen peroxide, Tea Tree Oil, purified water USP

Package Label - Principal Display Panel



Smallest Font Size: 7 Point



500 ml NDC: 81171-500-01

Sankind Hand Sanitizer 75 % v/v Isopropyl Alcohol

SANKIND HAND SANITIZER 75 % V/V ISOPROPYL ALCOHOL			
isopropyl alcohol gel			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:8 1171-500
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 mL in 100 mL
Inactive Ingredients			
Ingredient Name			Strength
GLYCERIN (UNII: PDC6A3C0OX)			
HYDROGEN PEROXIDE (UNII: BBX060AN9V)			
WATER (UNII: 059QF0KO0R)			
TEA TREE OIL (UNII: VIF565UC2G)			
HYPROMELLOSES (UNII: 3NXW29V3WO)			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:8 1171-500-01	500 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product	01/10/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	01/10/2021	

Labeler - UNIVET HOLDINGS LIMITED (988725024)**Registrant** - UNIVET HOLDINGS LIMITED (988725024)**Establishment**

Name	Address	ID/FEI	Business Operations
UNIVET HOLDINGS LIMITED		988725024	manufacture(8 1171-500)

Revised: 1/2021

UNIVET HOLDINGS LIMITED