SUAVE HAND SANITIZER PROFESSIONAL- alcohol gel
UNILEVER ASIA PRIVATE 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.

Suave Hand Sanitizer Professional - 75% Ethyl Alcohol

ACTIVE INGREDIENT

Ethyl alcohol 75% 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. Highly flammable liquid and vapour. Keep away from/do not spray near heat, hot surfaces, sparks, electrical appliances, open flames and other ignition sources. Do not store in car. No smoking. Keep containers tightly closed.

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, rinse cautiously with water for several minutes. Remove contact lenses if present and easy to do. Continue rinsing. If eye irritation persists: Get medical advice/attention.

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.

Keep out of reach of children. Causes serious eye irritation. If swallowed, get medical help or contact Poison Control Center right away.

DIRECTIONS

- Wet hands thoroughly with product and rub lightly until dry. Do not wipe off or rinse.

OTHER INFORMATION

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

INACTIVE INGREDIENTS

Acrylates Copolymer, Aminomethyl Propanol, Denatonium Benzoate, Glycerin, Purified Water USP, Tert-butyl Alcohol, Triethanolamine

SUAVE HAND SANITIZER PROFESSIONAL

alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:50069-110(NDC:73931-018)
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**

ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	75 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
BUTYL ACRYLATE/METHYL METHACRYLATE/METHACRYLIC ACID COPOLYMER (18000 MW) (UNII: JZ1374NL9E)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	
DENATONIUM BENZOATE (UNII: 4YK5Z54AT2)	
GLYCERIN (UNII: PDC6A3C0OX)	
WATER (UNII: 059QF0KO0R)	
TERT-BUTYL ALCOHOL (UNII: MD83SFE959)	
TROLAMINE (UNII: 9O3K93S3TK)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:50069-110-11	5000 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	08/25/2020	
2	NDC:50069-110-12	50 mL in 1 BOTTLE; Type 0: Not a Combination Product	08/25/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	08/25/2020	

Labeler - UNILEVER ASIA PRIVATE LIMITED (894632699)

Revised: 11/2020

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