SUAVE HAND SANITIZER PROFESSIONAL- alcohol gel
Guangdong Essence Daily Chemical Co., Ltd

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.

Drug Facts

Active ingredient

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.

Flammable, Keep away from heat or flame.

Do not use

- Onchildren 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 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.

Inactive ingredients

Purified Water USP, Acrylates Copolymer, Glycerin, Triethanolamine, Aminomethyl Propanol

other information

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

Packaging

SUAVE HAND SANITIZER PROFESSIONAL

alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:73931-017
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
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
TROLAMINE (UNII: 9O3K93S3TK)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	
DENATONIUM BENZOATE (UNII: 4YK5Z54AT2)	
TERT-BUTYL ALCOHOL (UNII: MD83SFE959)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73931-017-01	5000 mL in 1 BOTTLE; Type 0: Not a Combination Product	08/27/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	08/27/2020	

Labeler - Guangdong Essence Daily Chemical Co., Ltd (529796211)

Establishment

Name	Address	ID/FEI	Business Operations
Guangdong Essence Daily Chemical Co., Ltd		529796211	manufacture(73931-017)

Revised: 8/2020

Guangdong Essence Daily Chemical Co., Ltd