

I-SOFTTO INSTANT HAND SANITIZER- 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

Ethyl Alcohol 75% +/-5% (V/V)

Purpose

Antiseptic

Use

To help reduce bacteria on the skin and recommended for repeated use.

Warnings

For external use only.

Flammable,

Keep product away from fire or flame or sparks.

When using this product

do not use in or near eyes, In case of contact, rinse eyes thoroughly with water.

Stop use and ask a doctor if

irritation or rash appears on the skin.

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center right away.

Directions

- Place enough to cover both hands in the palm, and rub hands together until dry
- Children under 6 years of age should be supervised by an adult when applying this products.

Inactive ingredients

Water, Glycerin, Carbomer, Triethanolamine, Aloe Barbadensis Leaf Juice.

other information

- Do not store above 105 Fahrenheit.
- May discolor certain fabrics or surfaces.
- Harmful to wood finishes and plastics.

Packaging



i-Softto®

Instant Hand Sanitizer

Ethyl alcohol 75%±5% (V/V)

16.9 fl.oz / 500 ml

PREMIUM QUALITY
BY SOFTTO GROUP

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Other information • Do not store above 105°F • May discolor certain fabrics or surfaces • Harmful to wood finishes and plastics.

Inactive ingredients aloe barbadensis leaf juice, carbomer, glycerin, triethanolamine, water

Questions or comments? Call 86-020-81320487

Batch number/Expired Date: As shown on the package.
Manufacturer: Guangdong Essence Daily Chemical Co., Ltd
Distributed by: Guangzhou Softkiss cosmetics Co., Ltd
Address: No.88, Lane 2 of Softto street, Chong hua Economic and Technological Development Zone, Guangdong, China.

MADE IN CHINA

CONFORMITY PRODUCT Z01





I-SOFTTO INSTANT HAND SANITIZER

alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:73931-005
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)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73931-005-01	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/23/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC mono graph not final	part333E	06/23/2020	

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

Establishment

Name	Address	ID/FEI	Business Operations
Guangdong Essence Daily Chemical Co., Ltd		529796211	manufacture(73931-005)

Revised: 6/2020

Guangdong Essence Daily Chemical Co., Ltd