HAKU PIKAKE HAND SANITIZER- ethyl alcohol gel
SRC Company 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.

ACTIVE INGREDIENT

Ethyl alcohol 70%

INACTIVE INGREDIENTS

Water, Carbomer, Glycerin, Propylene Glycol, Triethanolamine, Polysorbate 80, Aloe barbadensis leaf extract, Fragrance

PURPOSE

ANTISEPTIC

WARNINGS

For external use only. Flammable. Keep away from heat or flame

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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.

Uses

• Hand sanitizer to help reduce bacteria that potentially can cause disease.
• Recommended for repeated use.

Directions

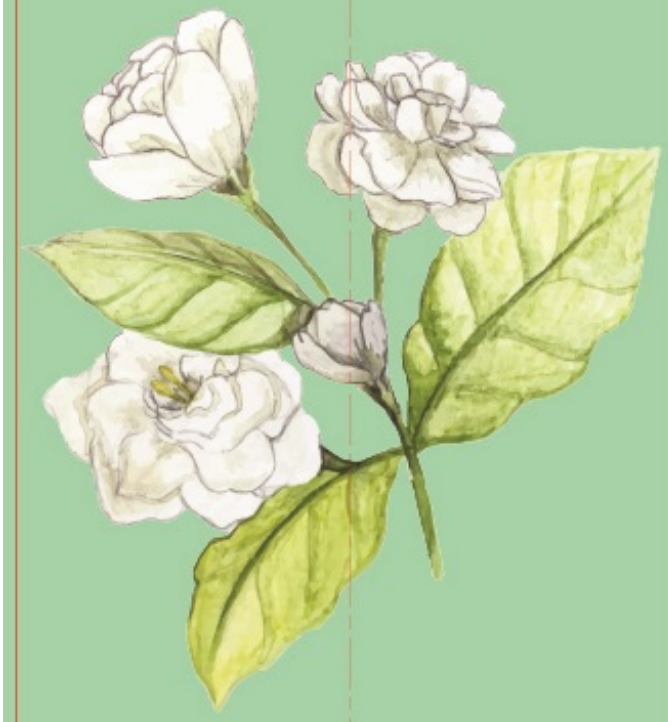
• Place enough product on hands to cover all surfaces. Rub hands together until dry.
• Supervise children under 6 years of age when using this product to avoid swallowing.
• Not recommended for infants

Other Information

• Store between 1□-30□ (33.8□-86□)
• Avoid freezing and excessive heat above 40□ (104□)

PACKAGE LABEL.PRINCIPAL DISPLAY PANEL

HAKU™



PIKAKE

HAND SANITIZER

2.02 fl. oz (60mL)

Drug Facts

Active ingredients — **Purpose**
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Inactive ingredients
Water, Carbomer, Glycerin, Propylene Glycol, Triethanolamine, Polysorbate 80, Aloe barbadensis leaf extract, Fragrance

Distributor:
SRC Company Co., Ltd.
15 Nonhyeon-ro 150-gil, Gangnam-gu, Seoul, Republic of Korea

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HAKU PIKAKE HAND SANITIZER

ethyl alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:78857-070
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	42 mL in 60 mL

Inactive Ingredients	
Ingredient Name	**Strength**
Water (UNII: 059QF0KO0R)	
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
Glycerin (UNII: PDC6A3C0OX)	
Propylene Glycol (UNII: 6DC9Q167V3)	
TROLAMINE (UNII: 9O3K93S3TK)	
Polysorbate 80 (UNII: 6OZP39ZG8H)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:78857-070-01	60 mL in 1 TUBE; Type 0: Not a Combination Product	06/01/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	06/01/2020	

Labeler - SRC Company Co., Ltd. (694702525)

Registrant - SRC Company Co., Ltd. (694702525)

Establishment

Name	Address	ID/FEI	Business Operations
C&N COSMETICS CO.,LTD		688853696	manufacture(78857-070)

Revised: 6/2020 SRC Company Co., Ltd.