NATURE HAND TOUCH SANITIZER GEL (ETHANOL)- alcohol liquid Dai Kyoung Pharmaceutic 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.

PLUS / Dai Kyoung Pharmaceutic Co.,Ltd. - Nature Hand Touch Sanitizer Gel (ethanol)

Alcohol

WATER, POLYSORBATE 20, CARBOMER, GLYCERIN

Hand sanitizer to help reduce bacteria that potentially can cause disease. For use when soap and water are not available.

keep out of reach of the children

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.

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 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.

for external use only

Drug Facts

Active Ingredient

Purpose

Alcohol 70 % ------ Antiseptic

Uses

■ Hand Sanitizer to help reduce bacteria that potentially can cause disease. For use when soap and water are not available.

Warnings

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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

water, polysorbate 20, carbomer, glycerin

NATURE HAND TOUCH SANITIZER GEL (ETHANOL)

alcohol liquid

D J		TC-		4º
Prod	HCT	into	rma	HON

Product TypeHUMAN OTC DRUGItem Code (Source)NDC:74279-0006

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	70 mL in 100 mL

Inactive Ingredients

mactive ingredients		
Ingredient Name	Strength	
WATER (UNII: 059QF0KO0R)		
POLYSORBATE 20 (UNII: 7T1F30V5YH)		
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0 A5MM307FC)		
GLYCERIN (UNII: PDC6A3C0OX)		

Packaging				
#	Item Code	Package Describtion		Marketing End Date
1	NDC:74279-0006- 1	100 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	04/10/2020	
Marketing Information				
	Marketing Categ	ry Application Number or Monograph Citation	Marketing Start Date	Marketing End Date

04/10/2020

Labeler - Dai Kyoung Pharmaceutic Co.,Ltd. (695045937)

OTC monograph not final part333A

Registrant - Dai Kyoung Pharmaceutic Co.,Ltd. (695045937)

Establishment			
Name	Address	ID/FEI	Business Operations
Dai Kyoung Pharmaceutic Co.,Ltd.		695045937	manufacture(74279-0006), label(74279-0006), pack(74279-0006)

Revised: 5/2020 Dai Kyoung Pharmaceutic Co.,Ltd.