

WONDER HAND GEL (ETHANOL)- alcohol gel

Dai Kyong 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 Kyong Pharmaceutic Co.,Ltd. - Wonder Hand Gel (Ethanol)

Alcohol

water, glycerin, tocopheryl acetate, triethanolamine, carbomer, polysorbate 20

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

Alcohol 72 %

Purpose

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, triethanolamine, tocopheryl acetate, glycerin, lemon flavor

WONDER HAND GEL (ETHANOL)

alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:74279-0009
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	540 mL in 750 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
CARBOMER HO MO POLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
GLYCERIN (UNII: PDC6A3C0OX)	
TROLAMINE (UNII: 9O3K93S3TK)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:74279-0009-1	750 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	04/10/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	04/10/2020	

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

Name	Address	ID/FEI	Business Operations
Dai Kyoung Pharmaceutic Co.,Ltd.		695045937	manufacture(74279-0009) , label(74279-0009) , pack(74279-0009)

Revised: 5/2020

Dai Kyoung Pharmaceutic Co.,Ltd.