CLEAN TO CLEAN HAND SANITIZER GEL PLUS- alcohol gel **ONA**

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.

ONA - ABETTERDAY PERFECT CLEAN HAND SANITIZER GEL

Alcohol

water, etc

Antiseptic

keep out of reach of the children

Place enough product in your palm to thoroughly spread on both hands and rub into the skin until dry.

Supervise children in the use of this product.

For external use only.

Flammable.

Keep away from fire or flame.

for external use only



Hand cleangel instant hand sanitizer





Drug Facts

Active Ingredient

Purpose

Ethyl Alcohol 70% V/V -

Antiseptic

Uses

- Hand sanitizer to help reduce bacteria on the skin.
- Recommended for repeated use.

Warnings

• For external use only. • Flammable. • Keep away from fire or flame

When using this product

- · avoid contact with eyes
- in case of eye contact immediately flush eyes with water, call a doctor
- avoid contact with broken skin.

Stop use and ask a doctor if

- Irritation and redness develops.
- condition persist for more than 72 hours.

Keep out of reach of children.

or contact a Poison Control Center right away.

Directions • Place enough product in your palm to thoroughly spread on both hands and rub into the skin until dry.

Supervise children in the use of this product.

Other Information

• Store at 1~30°C (34°F ~ 86°F) • May discolor some fabrics

Inactive Ingredients Water, Carbomer, Triethanolamine, Glycerin Propylene Glycol, Panthenol, Dipotassium Glycyrrhizate, Aloe Barbadensis Leaf Juice, Sodium Hyaluronate, Tocopheryl Acetate

Questions or Comments? email: Miglimkorea@gmail.com



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CLEAN TO CLEAN HAND SANITIZER GEL PLUS

alcohol gel

Product Information Product Type HUMAN OTC DRUG Item Code (Source) NDC:71060-070 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	
Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	

Packaging			
# Item Code Package Description		Marketing Start Date	Marketing End Date
1 NDC:71060-070- 01	110 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/09/2020	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	04/09/2020	

Labeler - ONA (689851847)

Registrant - ONA (689851847)

Establishment			
Name	Address	ID/FEI	Business Operations
KOREACOSPACK CO.,LTD.		689059789	manufacture(71060-070)

Establishment			
Name	Address	ID/FEI	Business Operations
ONA		689851847	label(71060-070), pack(71060-070)

Revised: 4/2020 ONA