

FIRST HAND SANITIZER- alcohol gel
Korea Life Science 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

ALCOHOL

INACTIVE INGREDIENT

glycerin carbomer, green tea extract, aloe extract, mugwort extract, L-menthol, triethanolamine

PURPOSE

SANITIZER

WARNINGS

1.Do not use the product on the following areas:

Around the eyes and ears, in the mouth, large body parts and damaged skin(Irritation may occur)

2.If you experience any of the following symptoms, discontinue use of the product immediately and consult a physician or pharmacist.

1) Hypersensitivity symptoms, e.g. rash, erythema, itchiness, and edema

2)Skin irritation symptoms

3.Other precautions

1) For external use only (do not swallow).

2) Avoid contact with eyes. If contact occurs, wash with clean water and consult a doctor or pharmacist.

3)Be careful not to inhale vapors in cases of extended or prolonged use. (Repeating inhaling of large amounts of ethanol vapor may cause irritation of the mucous membranes and headaches).

KEEP OUT OF REACH OF CHILDREN

Keep out of reach of children and go to the hospital immediately if swallowed.

Uses

Disinfection of hands and skin

Directions

Take an appropriate amount on your hands and rub thoroughly to dry.

PACKAGE LABEL.PRINCIPAL DISPLAY PANEL

FIRST HAND SANITIZER^(ETHANOL GEL)

Hand Sanitizer Gel

With Aloe & Green tea

Kills 99.9% of common germs*

 Contains : Ethanol 70% w/w

 Fragrance-free



500ml / 16.9 fl.oz.

FIRST HAND SANITIZER^(ETHANOL GEL) / 500ml

Drug Facts

Active Ingredients

Ethanol (Synthetic Alcohol) 70% ----- Disinfecting Agent *Purpose*

Uses

- instant healthcare personnel hand antiseptic to reduce bacteria that potentially can cause disease
- instant hand antiseptic to decrease bacteria on the skin

Warning

For external use only

When using this product

- if following abnormal symptoms persist, discontinue use
Irritation around the eyes, ears, mucous membranes, including the mouth, under the skin irritation and rashes
- Stop immediately and consult a doctor if you experience
 - 1) Hypersensitivity symptoms such as erythema, itching and dermatitis.
 - 2) Skin Irritation
 - 3) Following instructions when using medication
 - (1) For external use only (Do not use internally)
 - (2) Avoid getting into the eyes (if contact occurs, wash well with clean water)
- Be careful not to inhale or use excessively for a long time (ingesting ethanol repeatedly causes irritation to mucous membranes and headaches or other symptoms may appear. When used repeatedly in the same area, skin irritation may occur.
- Do not use the product for a long time in the same area as swelling, inflammation or sickness may occur due to absorption through the skin. It is not recommended to use this one areas that have been medically treated with a cast or bandage.
- Do not use in combination with soap or antibacterial cleansing agents.

Directions

- Pour a small amount into hands, spread evenly and rub into the skin

Other Information

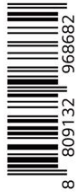
- read the directions and warnings before use
- avoid freezing and excessive heat above 40 degree C (104 degree F)

Inactive Ingredients

Water, Glycerin, Carbomer, Green Tea Extract, Aloe Extract(09), Mugwort Extract, L-Menthol, Triethanolamine

Questions? + 82 70 4237-3253

MADE IN KOREA



FIRST HAND SANITIZER

alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:74406-001
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	248 mL in 400 mL

Inactive Ingredients

Ingredient Name	Strength
GLYCERIN (UNII: PDC6A3C0OX)	
GREEN TEA LEAF (UNII: W2ZU1RY8B0)	
ARTEMISIA PRINCEPS LEAF (UNII: SY077EW02G)	
WATER (UNII: 059QF0KO0R)	
CARBOMER HOMO POLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
LEVOMENTHOL (UNII: BZ1R15MTK7)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:74406-001-01	500 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	05/19/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	05/19/2020	

Labeler - Korea Life Science Co.,ltd (694914835)

Registrant - Korea Life Science Co.,ltd (694914835)

Establishment

Name	Address	ID/FEI	Business Operations
Korea Life Science Co.,ltd		694914835	manufacture(74406-001)

Revised: 5/2020

Korea Life Science Co.,ltd