HAEA PLUS HAND SANITIZER- alcohol gel Dr.s Medi 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.

Dr.s Medi Co., Ltd. - Haea plus Hand Sanitizer

Alcohol

water, chitosan (aqueous solution 1%), glycerin, allantoin, propolis extract, fragrance

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















Useful in public places! (hospitals, toilets, public transportation, etc.)

[Product Name] Hae · a plus Hand Sanitizer

[Active Ingredient] Ethanol 70%

[Capacity] 17ml

[Other Additives] Chitosan (aqueous solution 1%), Glycerin, allantoin, purified water, propolis extract, fragrance rose

[Efficacy] Sterilization of hands and skin

[Usage and Dose]

Spray or put an appropriate amount on your hands, rub and dry. [Precautions for Use]

- 1. Do not use on the following body parts.
- In your mouth, on mucous membranes or damaged skin (it may irritate.) 2. Stop use immediately and consult a dermatologist if:
- 1) Hypersensitivity symptoms such as rash or itching occur Skin irritation symptoms occur
 Other use precautions
 I) It is external use only.
- 2) Be careful not to get it into your eyes, and if so, rinse thoroughly with water.
- 3) Be careful not to inhale vapor when using it extensively or for a long time. If you inhale ethanol vapor in large quantities or repeatedly, irritation to the mucous membrane or headache may occur.
- 4. Storage precautions
- 1) Keep it away from places with high temperatures or fire.
- 2) Keep it out of the reach of children. If a child swallows it, go to the hospital
- 3) Taking it out of the original container and storing it in another container may cause a misuse incident or quality deterioration. Put it in the original container and keep it tightly closed.

[Expiration Date and Manufacturing Number] Separately indicated at the bottom of the container

[Storage] Airtight container, store at room temperature (1 to 30 °C)

[R&D/Technology Support Distributor] SUN Life Science Co.,Ltd. 44-1, Geumbit-ro, Paju-si, Gyeonggi-do, Republic of Korea

[Manufacturer] AseaPharm Co., Ltd.

65, Yongdu-ro 18beon-gil, Daegotmyeon, Gimpo-si, Gyeonggi-do, Republic of Korea [Distributor] Dr's MEDI Co., Ltd.

Suite 804, Kintex Plaza, 1573, Jungang-ro, Ilsanseo-gu, Goyang-si, Gyeonggi-do, Republic of Korea

[Consumer Help Desk] +82-31-922-2240

* This product may be compensated according to the Fair Trade Commission Notice.

Quasi Drug



HAEA PLUS HAND SANITIZER

alcohol gel

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:76670-0001

TOPICAL Route of Administration

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	11.9 mL in 17 mL

Inactive Ingredients

Ingredient Name	Strength
Ingredient rume	Strength

WATER (UNII: 059QF0KO0R)

GLYCERIN (UNII: PDC6A3C0OX)	
PROPOLIS WAX (UNII: 6 Y8 XYV2NOF)	
ALLANTO IN (UNII: 344S277G0Z)	

	Packaging			
;	# Item Code	Package Description	Marketing Start Date	Marketing End Date
	NDC:76670-0001-	17 mL in 1 BOTTLE, SPRAY; 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 - Dr.s Medi Co.,Ltd. (694505169)

Registrant - Dr.s Medi Co.,Ltd. (694505169)

Establishment			
Name	Address	ID/FEI	Business Operations
Dr.s Medi Co.,Ltd.		694505169	label(76670-0001), manufacture(76670-0001)

Revised: 4/2020 Dr.s Medi Co.,Ltd.