

**DR. GREENDAY S HAND SANITIZER- alcohol liquid
WOWBIOTECH**

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.

WOWBIOTECH - DR. GREENDAY S HAND SANITIZER

Alcohol

water, triethanolamine, camellia sinensis leaf extract, glycerin, carbomer, propylene glycol

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

Dr. Greenday S
Hand Sanitizer

닥터 그린데이 에스 손소독제

hand sanitizer used
without water
99.9% sanitization

100ml / 3.38 fl. oz
MADE IN KOREA

Dr. Greenday S Hand Sanitizer

Drug Facts

Active Ingredient	Purpose
Ethanol 70 %	Sanitizers

Uses

- Disinfection of hands and skin

Warnings

- Do not use on the following body parts. Around the eyes and ears, in the oral cavity, a wide range of body parts and damaged skin (may have irritating effects)
- If the following symptoms appear, stop using them immediately and consult a doctor or pharmacist. 1)Irrpersensitive symptoms such as rash, erythema, itching, edema 2)Skin irritation symptoms
- Other precaution
1)For external use only 2)Be careful not to get into your eyes, and if so, rinse well, clean water and consult a doctor or pharmacist. 3)Be careful do not irritate vapor when using it extensively or for a long time. (if you drink ethanol vapor in large quantity or repeatedly, irritation to the mucous membrane, headache, etc., may occur. (only for products containing ethanol) 4)When repeated use on the same skin, be careful as the skin may become rough due to degreasing. 5)When used in sealed bandages, cast bandages, packs, etc., irritation symptoms may appear. 6)Do not use this medicine for appear anal or vaginal poultice as it may cause irritation or chemical burns. 7)Do not use for any purpose.

Precautions for storage

- Avoid shading and keep in shading.
- Keep it out of the reach of children, and if a child swallows it, go to the hospital right away.
- After use, close the product completely with a lid to prevent the product from drying out or entering foreign objects.
- Taking it out of the original container and storing it in another container may cause accidents due to misuse or deterioration in quality, so store it in the original container.

Directions

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

Other information

- How to save
Airtight container, storage at room temperature. (1-30°C)

Inactive Ingredients	
<ul style="list-style-type: none"> Ethanol, Water, Triethanolamine, Camellia sinensis leaf extract, Glycerin, Carbomer, Propylene Glycol 	

Manufactured By WOWBIOTECH
30, 67beon-gil Gajeong-ro Seo-gu, Incheon, Republic of KOREA

DR. GREENDAY S HAND SANITIZER

alcohol liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:76627-0011
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	350 mL in 500 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
TROLAMINE (UNII: 9O3K93S3TK)	
CARBOMER HO MO POLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
GLYCERIN (UNII: PDC6A3C0OX)	

GREEN TEA LEAF (UNII: W2ZU1RY8B0)

BUTYLENE GLYCOL (UNII: 3XUS85K0RA)

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:76627-0011-1	500 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	05/25/2020	
2	NDC:76627-0011-2	500 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	05/25/2020	05/29/2020

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	05/25/2020	

Labeler - WOWBIOTECH (695625834)

Registrant - WOWBIOTECH (695625834)

Establishment

Name	Address	ID/FEI	Business Operations
WOWBIOTECH		695625834	manufacture(76627-0011) , label(76627-0011) , pack(76627-0011)

Revised: 6/2020

WOWBIOTECH