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

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 HOMOPOLYMER, 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