

GERMS BE GONE HAND SANITIZER- alcohol liquid

Brands International

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.

Germes Be Gone Hand sanitizer

Drug Facts

Active Ingredient

Ethyl Alcohol, 75% v/v

Purpose

Antiseptic (skin) clenser

Uses

- Antiseptic (skin) clenser

Warnings

For external use only. Flammable, keep away from fire or flame.

When using this product keep out of eyes. Avoid contact with eyes. If contact with eyes occurs, rinse thoroughly with water.

Stop use and ask a doctor if irritation develops.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions

- For occasional and personal domestic use.
- Rub thoroughly into hands for at least 30 seconds. Allow to dry.
- Supervise children in the use of this product.

Other Information

- Store below 43°C (110°F)
- may discolor certain fabrics or surfaces

Inactive Ingredients

carbomer, aloe barbadensis leaf juice, aminomethyl propanol, glycerin, propylene glycol,

water, isopropyl myristate, tocopheryl acetate (Vit E).

Please see label on the container.



GERMS BE GONE HAND SANITIZER

alcohol liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:50157-520
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	75 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
WATER (UNII: 059QF0KO0R)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED) (UNII: 59TL3WG5CO)	
GLYCERIN (UNII: PDC6A3C0OX)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:50157-520-59	59 mL in 1 BOTTLE; Type 0: Not a Combination Product	12/03/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	12/03/2020	

Labeler - Brands International (243748238)

Registrant - Brands International (243748238)

Establishment

Name	Address	ID/FEI	Business Operations
Brands International		243748238	manufacture(50157-520)

Revised: 4/2021

Brands International