GERM END ANTIBACTERIAL HAND SANITIZER- ethyl alcohol gel
Lee's Collection Inc.

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.

GermEnd ANTIBACTERIAL HAND SANITIZER

Drug Facts

Active ingredient

Ethyl alcohol 75%

Purpose :

Antiseptic

Use ▪ to decrease bacteria on the skin that could cause disease ▪

Warning

For external use only : Hands.

Flammable: keep away from fire and flame.

Use with caution for people with alcohol allergies.

When using this product ▪ Keep out of eyes. In case of contact with eyes, flush thoroughly with water ▪ Avoid contact with broken skin. Do not inhale or ingest.

Stop use and ask a doctor if irritation and redness develop and condition persists for more than 72 hours.

Keep out of reach of children.

Directions Wet hands thoroughly with product and allow to dry without wiping. For children under 6, use only under adult supervision. Not recommended for infants.

Other information ▪ Store between 15-30C (59-86F) ▪ Avoid freezing and excessive heat above 40C (104F)

Inactive ingredients: water, glycerin, propylene glycol, carbo-mer, aloe vera extract, etc

Kills 99.99% of germs

Effective at eliminating over 99.99% of common harmful germs and bacteria in less than 30 seconds.

Distributed by ▪ Lee's Collection Inc.

S.EI Monte ,CA 91733

PD:04/16/2020

EXP:04/15/2022

Made in China

Packaging

GERM END ANTIBACTERIAL HAND SANITIZER

ethyl alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:74027-200
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
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:74027-200-06	60 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/20/2020	
2	NDC:74027-200-07	80 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/20/2020	
3	NDC:74027-200-08	100 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/20/2020	
4	NDC:74027-200-09	250 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/20/2020	
5	NDC:74027-200-10	500 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/20/2020	
6	NDC:74027-200-11	750 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/20/2020	
7	NDC:74027-200-12	1000 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/20/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	04/20/2020	

Labeler - Lee's Collection Inc. (022215540)

Revised: 4/2020 Lee's Collection Inc.