

GUARD HAND SANITIZER- ethyl alcohol gel
Apollo Health and Beauty Care 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.

Drug Facts

Active Ingredients

Ethyl Alcohol 65%

Purpose

Antiseptic

Uses

For Personal hand hygiene to help the spread of bacteria and can be used in place of hand washing if soap and water are not available.

Warnings

For External use only

- Flammable
- Keep away from source of heat or fire.

When using this product

Avoid contact with eyes. If contact occurs, rinse thoroughly with water. Do not Inhale.

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Stop use and sak a doctor

If irritation or redness develops and lasts

Keep out of reach of children

In case of accidental ingestion, get medical help or contact a Poison Control Centre immedoately

Directions

- Adults only
- For occasional and personal use
- Rub thoroughly into hands for at least 30 seconds. Allow to dry

Other information

- Store at a temperature below 110⁰F (43⁰C)
- May discolr certain fabrics or surfaces

Inactive Ingredients

Water, PEG-6, Fragrance, Isopropyl Alcohol, Glycerin, Isopropyl Myristate, Aloe Barbadensis Leaf Juice, Tocopheryl Acetate, AMP-Acrylates/Vinyl Isodecanate Crosspolymer

Questions or comments ?

1-866-695-3030

Label copy



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Warnings For external use only. ■ flammable ■ keep away from source of heat or fire

When using this product avoid contact with eyes. If contact occurs, rinse thoroughly with water. Do not inhale.

Do not use ■ on children less than 2 months of age ■ on open skin wounds ■ on broken or damaged skin ■ if you are pregnant or breastfeeding

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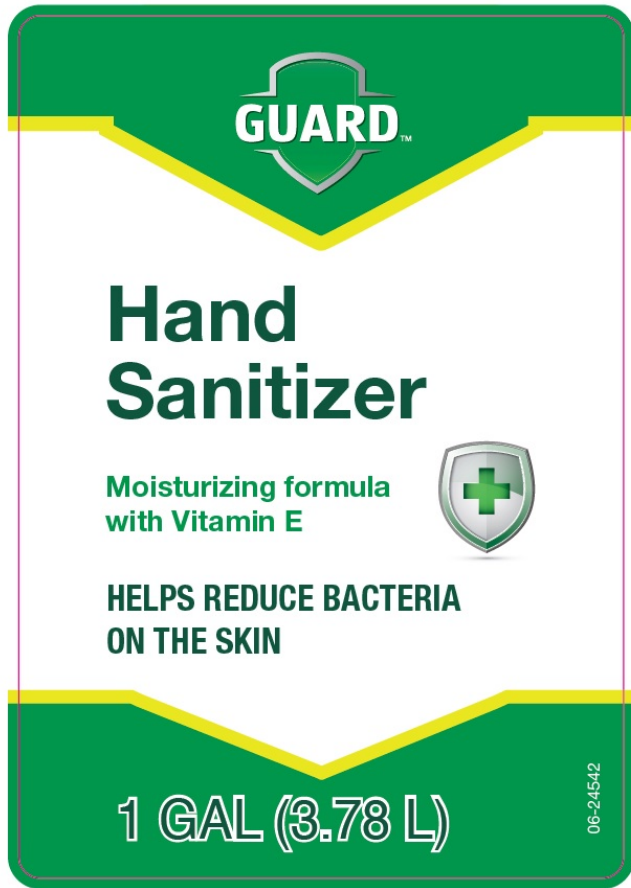
Questions or comments? 1-866-695-3030

Distributed By: Apollo Health and Beauty Care Inc.
1 Apollo Place, Toronto, ON, Canada, M3J 0H2

Made in Canada with domestic and imported components.



06-24543



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GUARD HAND SANITIZER

ethyl alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63148-517
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	650 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0K00R)	
POLYETHYLENE GLYCOL 300 (UNII: 5655G9Y8AQ)	
ACRYLATES/VINYL ISODECANOATE CROSSPOLYMER (10000 MPAS NEUTRALIZED AT 0.5%) (UNII: 2N8MDB79NA)	
GLYCERIN (UNII: PDC6A3C0OX)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
.ALPHA.-TOCOPHEROL ACETATE, DL- (UNII: WR1WPI7EW8)	

ALOE VERA LEAF (UNII: ZY81Z83H0X)

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63148-517-09	3780 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/06/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	10/06/2020	

Labeler - Apollo Health and Beauty Care Inc (201901209)

Registrant - Apollo Health and Beauty Care Inc (201901209)

Establishment

Name	Address	ID/FEI	Business Operations
Apollo Health and Beauty Care Inc		201901209	manufacture(63148-517)

Revised: 10/2020

Apollo Health and Beauty Care Inc