



COVID-19 is an emerging, rapidly evolving situation.

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Get the latest public health information from CDC: <https://www.coronavirus.gov>

Get the latest research information from NIH: <https://covid19.nih.gov>

Learn more about COVID-19 and you from HHS: <https://combatcovid.hhs.gov>



LABEL: TOPCARE HAND SANITIZER- ethyl alcohol gel

VIEW PACKAGE PHOTOS



Contains inactivated NDC Code(s)

NDC Code(s): 36800-235-01

Packager: TOPCO ASSOCIATES LLC

Category: HUMAN OTC DRUG LABEL

DEA Schedule: None

Marketing Status: OTC monograph not final

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 LABEL INFORMATION

Updated November 18, 2011

If you are a consumer or patient please visit [this version](#).

CLOSE ALL SECTIONS

ACTIVE INGREDIENT

Ethyl Alcohol 62%

CLOSE

PURPOSE

Antimicrobial

CLOSE

USES

To help reduce bacteria on the skin that could cause disease. Recommended for repeated use.

CLOSE

WARNINGS

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

When using this product

Avoid contact with eyes. If contact occurs, rinse with water.

Stop using this product and ask doctor if

Irritation or rash develops and lasts.

Keep out of reach of children.

In case of accidental ingestion, get medical help or contact Poison Control Center immediately.

CLOSE

DIRECTIONS

Spray enough product in your palm to thoroughly cover your hands. Rub hands together briskly until dry. Children under 6 years of age should be supervised when using this product.

CLOSE

OTHER INFORMATION

Store below 110 F (43 C)

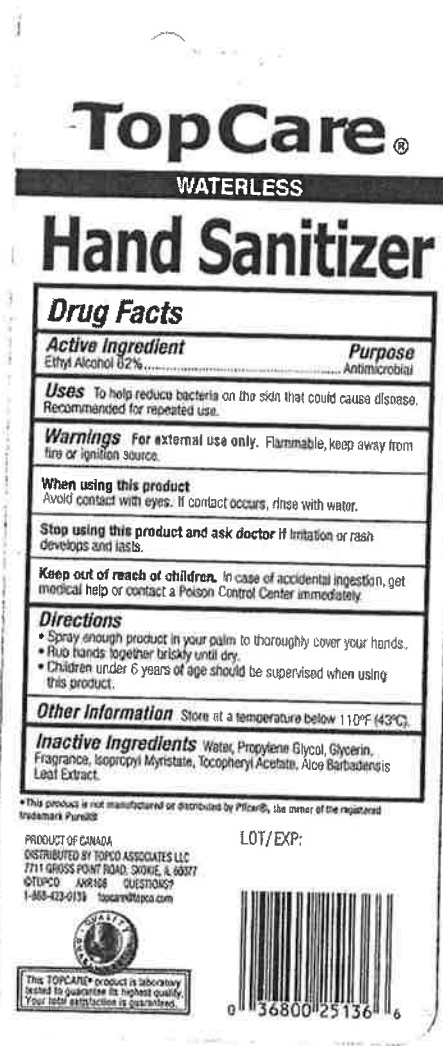
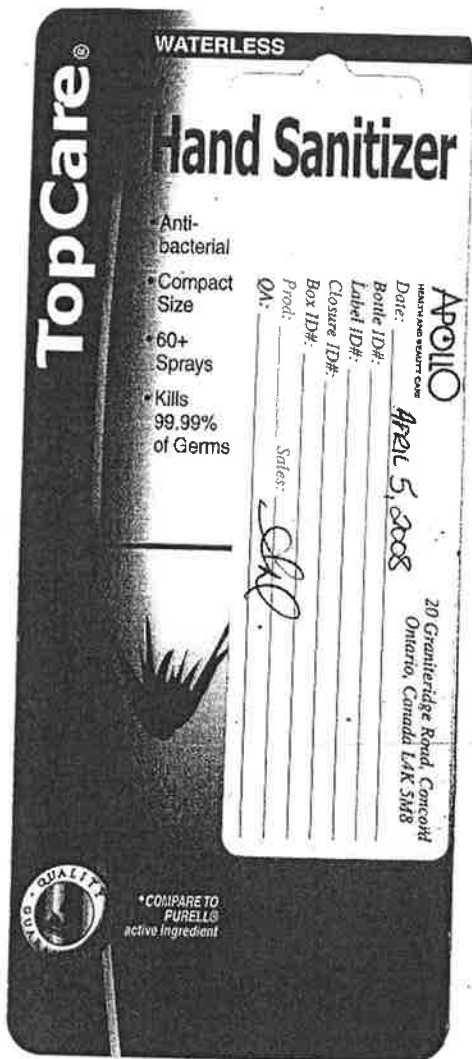
CLOSE

INACTIVE INGREDIENTS

Water, Propylene Glycol, Glycerin, Fragrance, Isopropyl Myristate, Tocopheryl acetate, Aloe Barbadosis Leaf Extract.

CLOSE

LABEL COPY



CLOSE

INGREDIENTS AND APPEARANCE

TOPCARE HAND SANITIZER ethyl alcohol gel				
PRODUCT INFORMATION				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:36800-235	
Route of Administration	TOPICAL			
ACTIVE INGREDIENT/ACTIVE MOIETY				
Ingredient Name		Basis of Strength	Strength	
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)		ALCOHOL	62 mL in 100 mL	
INACTIVE INGREDIENTS				
Ingredient Name			Strength	
WATER (UNII: 059QF0K00R)				
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)				
GLYCERIN (UNII: PDC6A3C0OX)				
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)				
TOCOPHEROL (UNII: R0ZB2556P8)				
ALOE VERA LEAF (UNII: ZY81Z83H0X)				
PACKAGING				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:36800-235-01	7.4 mL in 1 BOTTLE, SPRAY		
MARKETING INFORMATION				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part333E	11/20/2011		
LABELER - TOPCO ASSOCIATES LLC (006935977)				
REGISTRANT - APOLLO HEALTH AND BEAUTY CARE (201901209)				
ESTABLISHMENT				
NAME	ADDRESS	ID/FEI	BUSINESS OPERATIONS	

APOLLO HEALTH AND BEAUTY CARE	201901209	manufacture
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[CLOSE](#)

[CLOSE ALL SECTIONS](#)

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SAFETY

[Report Adverse Events](#), [FDA Safety Recalls](#), [Presence in Breast Milk](#)

RELATED RESOURCES

[Medline Plus](#), [Clinical Trials](#), [PubMed](#), [Biochemical Data Summary](#)

MORE INFO ON THIS DRUG

[View Labeling Archives](#), [RxNorm](#), [Get Label RSS Feed](#), [View NDC Code\(s\)](#)NEW!

