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)

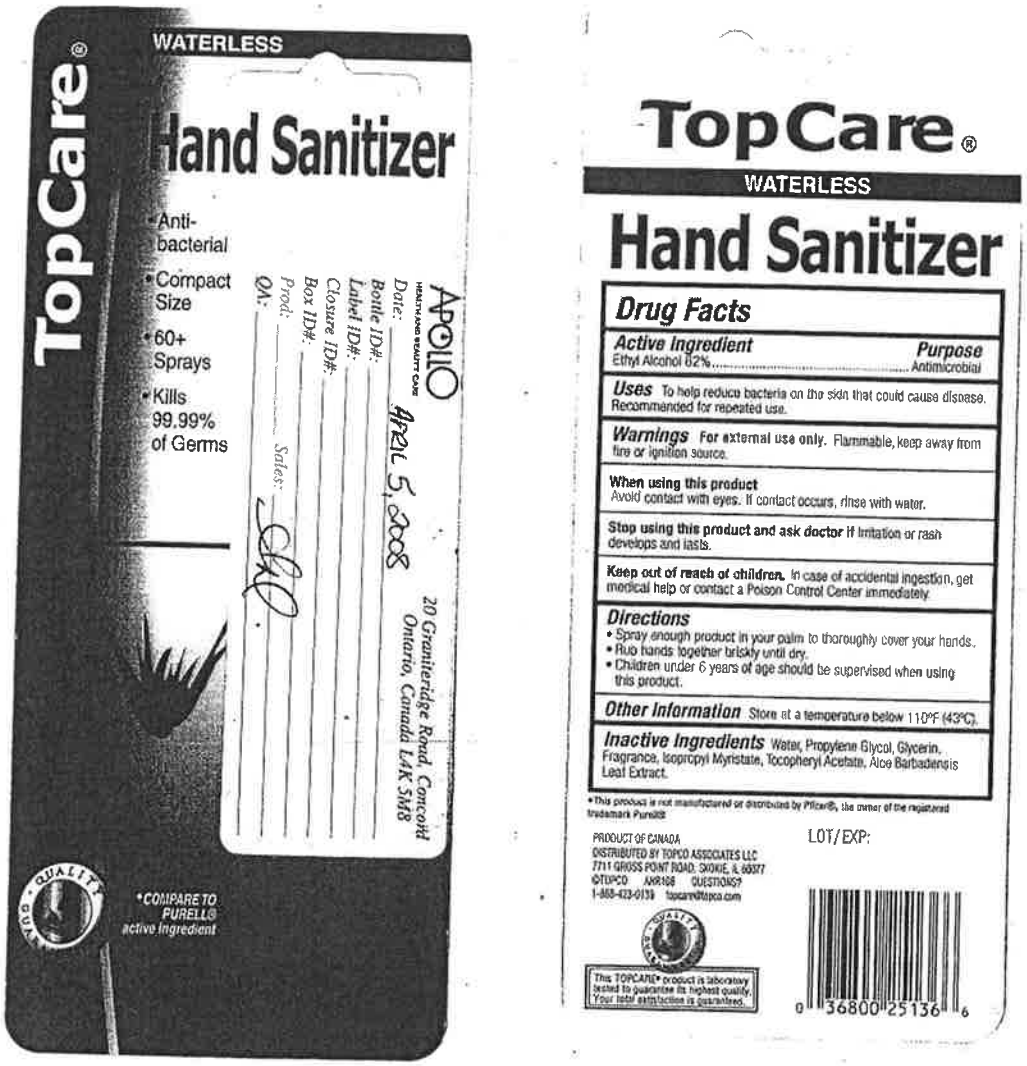
CLOSE

INACTIVE INGREDIENTS

Water, Propylene Glycol, Glycerin, Fragrance, Isopropyl Myristate, Tocopheryl acetate, Aloe Barbadensis 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: 059QF0KO0R)	
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

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!