

HAND SANITIZER WITH MOISTURIZERS AND VITAMIN E- alcohol liquid

Family Dollar Services 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 Ingredient

□Ethyl Alcohol 65%

Purpose

Antiseptic

Uses • hand sanitizer to help reduce bacteria on the skin. • recommended for repeated use

Warnings • Flammable • Keep away from fire or flame • For external use only

When using this product • avoid contact with eyes • in case of eye contact immediately flush eyes with water, call a doctor • avoid contact with broken skin

Discontinue use if irritation or redness develops. If condition persists for more than 72 hours, consult a doctor.

Keep out of reach of children. Children should only use this product under adult supervision.

Do not drink. Not edible. In case of accidental ingestion, seek professional assistance or contact a Poison Control Center right away.

Other Information •do not store above 105F. May discolor some fabrics • Harmful to wood finishes and plastics

Directions •place enough product in your palm to thoroughly spread on both hands and rub into the skin until dry • recommended for repeated use.

Inactive ingredients aloe barbadensis leaf extract, caprylyl glycol, carbomer, dimethicone, fragrance, phenoxyethanol, propylene glycol, tocopheryl acetate, triethanolamine, water.



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*This product is not manufactured or distributed by G10 Industries, Inc., distributor of Purell®	

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alcohol liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:55319-505
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	65 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	

PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
TROLAMINE (UNII: 9O3K93S3TK)	
WATER (UNII: 059QF0K00R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:55319-505-34	10 mL in 1 BOTTLE, SPRAY		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	10/01/2012	

Labeler - Family Dollar Services Inc. (024472631)

Revised: 9/2012

Family Dollar Services Inc.