

HAND SANITIZER- ethyl alcohol gel
DOLGENCORP, LLC

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

Active ingredient

Ethyl alcohol 65%

Purpose

Purpose

Antisepetic

Use

Use

- to decrease bacteria on the skin that could cause disease
- recommended for repeated use

Warnings

Warnings

For external use only-hands

flammable

Flammable. Keep away from heat and flame

When using this product

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

Stop use and ask a doctor if skin irritation develops

Keep out of reach of children

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions

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

Inactive ingredients

Inactive ingredients aloe barbadensis leaf juice, benzophenone-4, blue 1, carbomer, fragrance, glycerin, isopropyl myristate, propylene glycol, red 33, tocopheryl acetate, water

other information**Other information:**

- Do not store above 105°F.
- May discolor some fabrics
- Harmful to wood finishes and plastics

claims

Effective at eliminating 99.99% of many common harmful germs and bacteria in as little as 15 seconds

MADE IN U.S.A. WITH U.S. AND FOREIGN COMPONENTS

Adverse reactions section

DISTRIBUTED BY DOLGENCORP, LLC

100 MISSION RIDGE, GOODLETTSVILLE, TN 37072

Principal Display Panel

DG health

Instant

Hand

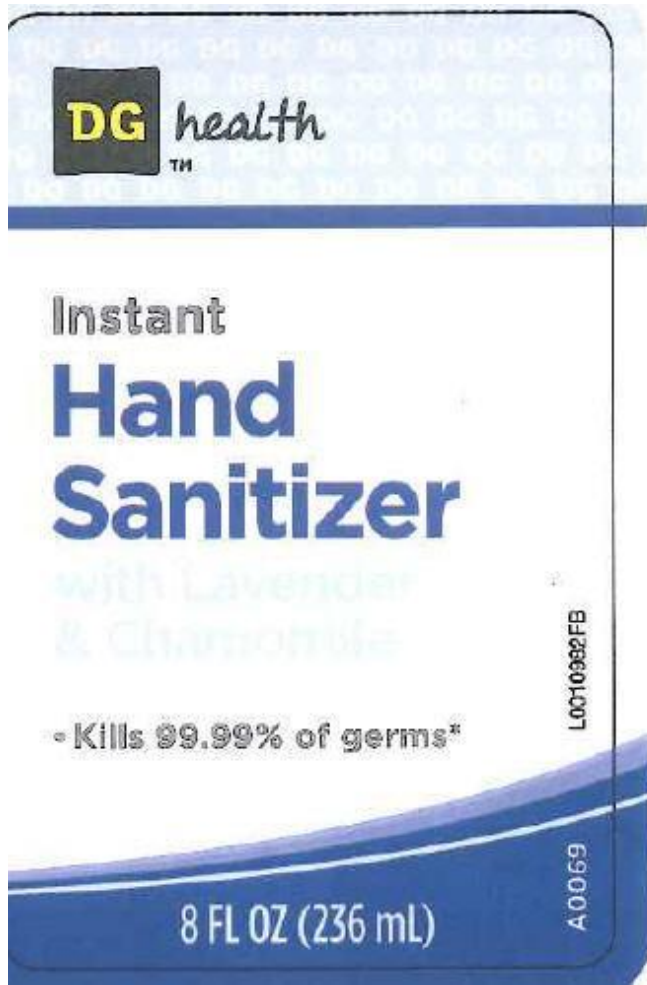
Sanitizer

with Lavender

+ Chamomile

- Kills 99.99% of germs

8 FL OZ (236 mL)



Drug Facts	
Active Ingredient Ethyl Alcohol 65%	Purpose Antiseptic
Use ■ to decrease bacteria on the skin that could cause disease ■ recommended for repeated use	
Warnings For external use only-hands Flammable. Keep away from heat and flame. When using this product ■ keep out of eyes. In case of contact with eyes, flush thoroughly with water. ■ do not inhale or ingest ■ avoid contact with broken skin Stop use and ask a doctor if skin irritation develops. Keep out of reach of children. In case of accidental ingestion, seek professional assistance or contact a Poison Control Center immediately.	
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 ■ do not store above 105°F ■ may discolor some fabrics ■ harmful to wood finishes and plastics	
Inactive Ingredients aloe barbadensis leaf juice, benzophenone-4, blue 1, carbomer, fragrance, glycerin, isopropyl myristate, propylene glycol, red 33, tocopheryl acetate, water	
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100% QUALITY GUARANTEED (888) 309-9030 DISTRIBUTED BY DOLGENCORP, LLC 100 MISSION RIDGE, GOODLETTSVILLE, TN 37072 MADE IN U.S.A. WITH U.S. AND FOREIGN COMPONENTS	



HAND SANITIZER

ethyl alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:559 10-523
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

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

Inactive Ingredients

Ingredient Name	Strength
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
SULISOBENZONE (UNII: 1W6L629B4K)	
FD&C BLUE NO. 1 (UNII: HBR47K3TBD)	
CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED) (UNII: 59TL3WG5CO)	
GLYCERIN (UNII: PDC6A3C0OX)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	

D&C RED NO. 33 (UNII: 9DBA0SBB0L)

.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)

WATER (UNII: 059QF0K00R)

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:55910-523-34	236 mL in 1 BOTTLE, PUMP		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	08/19/2010	

Labeler - DOLGENCORP, LLC (068331990)

Registrant - Vi-Jon (790752542)

Establishment

Name	Address	ID/FEI	Business Operations
Vi-Jon		088520668	manufacture(55910-523)

Revised: 11/2012

DOLGENCORP, LLC