## HAND SANITIZER- 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.

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#### **Drug Facts**

#### **Active ingredients**

#### **Active ingredient**

Ethyl alcohol 65%

#### **Purpose**

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Antispetic

#### 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 abut 105° IF.
- May discolor sme fabrics
- Harmful to wood finishes and plastics

#### claims

Effective at eliminating 99.99% of many common harmful germs and bacteria in as littlw as 15 seconds MADE IN U.S.A. WITH U.S. AND FOREIGN COMPONENTS

#### Adverse reactions section

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#### **Principal Display Panel**

DG health

Instant

Hand

Sanitizer

with Lavender

- + Chamomile
- Kills 99.99% of germs

8 FL OZ (236 mL)





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

L0010982BC

#### HAND SANITIZER

ethyl alcohol gel

# Product Information Product Type HUMAN OTC DRUG LABEL Item Code (Source) NDC:55910-523 Route of Administration TOPICAL DEA Schedule

## Active Ingredient/Active Moiety Ingredient Name Basis of Strength ALCOHOL (ALCOHOL) ALCOHOL ALCOHOL 650 mL in 1 mL

Inactive Ingredients		
Ingredient Name	Strength	
ALOE VERA LEAF		
SULISOBENZONE		
FD&C BLUE NO. 1		
CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED)		
GLYCERIN		
ISOPROPYL MYRISTATE		
PROPYLENE GLYCOL		

D&C RED NO. 33	
.ALPHATO COPHEROL ACETATE	
WATER	

1	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