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.

Hand Sanitzer
523

Active ingredient

Ethyl alcohol 65%

Purpose

Antispetic

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
- avoid contact with broken skin
- do not inhale or ingest

Stop use and ask a doctor if

skin irritation develops

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center right away.

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

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

Other information

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

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

100% QUALITY GUARANTEED (888-309-9030)

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

DISTRIBUTED BY DOLGENCORP, LLC

100 MISSION RIDGE, GOODLETTSVILLE, TN 37072

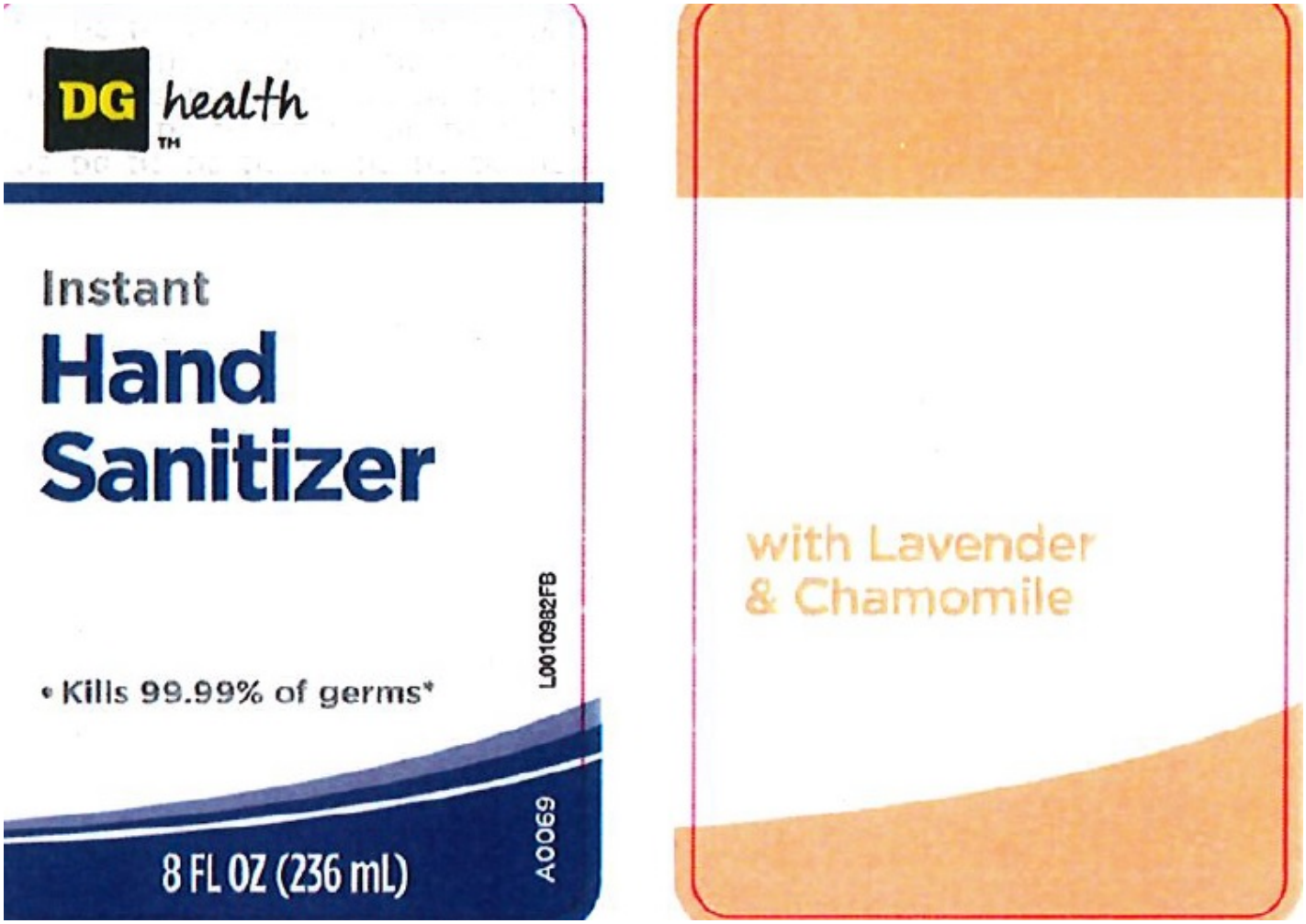
523.000/523AA

principal display panel

DG health Instant Hand Sanitizer with Lavender & Chamomile

- Kills 99.99 of germs

8 FL OZ (236 mL)



HAND SANITIZER ethyl alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:55910-523
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: H3R47K3TBD)	
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: 059QF0KO0R)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:55910-523-34	236 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	08/19/2010	

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: 5/2020 DOLGENCORP, LLC