ETHYL ALCOHOL- ethyl alcohol liquid
DOLLAR GENERAL CORPORATION

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.

Moisturizing Hand Sanitizer
849.001/849AA-AG rev 0, rev 1 & 849BA

Active ingredient

Ethyl Alcohol 62%

Purpose

Antiseptic

Uses

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

Warnings

For exterenal use only: hands

Flammable, keep away from fire or flame.

When using this product

- keep out of eyes. In case of contact with eyes, flush thoroghly with water.
- avoid contact with broken skin
- do not inhale or ingest

Stop use and ask a doctor if

- irritation and redness develop
- condition persists for more than 72 hours

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

Other information

- do not store above 105ºF
- may discolor some fabrics
- harmful to wood finishes and plastics

Inactive ingredients

water, carbomer or acrylates/C10-30 alkyl acrylate crosspolymer, fragrance, glycerin, isopropyl myristate, tocopheryl acetate

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

100% Satisfaction Guaranteed! (888)309-9030

DISTRIBUTED BY OLD EAST MAIN CO.

100 MISSION RIDGE

GOODLETTSVILLE, TN 37072

Principal display panel

STUDIO SELECTION

moisturizing

HAND SANITIZER

original

- With Vitamin E
- kills 99.99% of germs & bacteria*

8 FL OZ (236 mL)

Principal display panel

Since 1903

Rexall

Moisturizing

Hand Sanitizer

Kills 99.99% of Germs*

enriched with moisturizers

32 FL. OZ. (1QT) (946 mL)

ETHYL ALCOHOL

ethyl alcohol liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:75712-849
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)	
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED) (UNII: 59TL3WG5CO)	

GLYCERIN (UNII: PDC6A3C0OX)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
TOCOPHEROL (UNII: R0ZB2556P8)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:75712-849-34	236 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/01/2023	
2	NDC:75712-849-16	59 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/01/2023	
3	NDC:75712-849-45	946 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/01/2023	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	05/01/2023	

Labeler - DOLLAR GENERAL CORPORATION (006946172)

Registrant - Vi Jon, LLC (790752542)

Establishment

Name	Address	ID/FEI	Business Operations
Vi Jon, LLC		088520668	manufacture(75712-849)

Establishment

Name	Address	ID/FEI	Business Operations
Vi Jon, LLC		790752542	manufacture(75712-849)

Revised: 5/2023 DOLLAR GENERAL CORPORATION