

LUCKY ALCOHOL FREE HAND SANITIZER VITAMIN E- benzalkonium chloride liquid
Delta Brands, 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.

Lucky Alcohol Free hand sanitizer-Vit E

Active Ingredient

Benzalkonium Chloride 0.13%

Purpose

Antiseptic

Uses

hand sanitizer to help reduce bacteria on the skin that could cause disease

Warnings

For external use only

When using this product

do not get into the eyes. If contact occurs, rinse eyes thoroughly with water.

Stop use and ask a doctor if

if irritation or rash appears and lasts

Keep out of reach of children

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

Directions

■ place enough product in your palm ■ rub hands vigorously until dry ■ children under 6 years of age should be supervised when using this product

Other information

■ store below 106°F (41°C) ■ may discolor certain fabrics or surfaces

Inactive ingredients

water, glycerin, carbomer, poly sorbate 20, PEG-40 hydrogenated castor oil, fragrance, sodium hydroxide, methylchloroisothiazolinone, methylisothiazolinone, tocopheryl acetate

Package Label



ALCOHOL-FREE



instant
hand sanitizer
Vitamin E

11702/01-F(001)



Vitamin E

Kills 99.9%
of Germs*

8FLOZ (236mL)

Drug Facts

Active ingredient	Purpose
Benzalkonium chloride 0.13%	Antiseptic

Use hand sanitizer to help reduce bacteria on the skin that could cause disease

Warnings
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Inactive ingredients water, glycerin, carbomer, polysorbate 20, PEG-40 hydrogenated castor oil, fragrance, sodium hydroxide, methylchloroisothiazolinone, methylisothiazolinone, tocopheryl acetate

*99.9% of most common germs.

ITEM NO: 11702
11702/01-R(001)

Manufactured for
DELTA BRANDS INC.
1890 Palmer Avenue
Larchmont, New York 10538 USA
www.deltabrands.com
MADE IN TURKEY



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LUCKY ALCOHOL FREE HAND SANITIZER VITAMIN E

benzalkonium chloride liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:20276-139
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.13 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength

WATER (UNII: 059QF0K00R)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
GLYCERIN (UNII: PDC6A3C0OX)	
POLYOXYL 40 HYDROGENATED CASTOR OIL (UNII: 7YC686GQ8F)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:20276-139-08	236 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/10/2020	
2	NDC:20276-139-10	296 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/10/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	03/10/2020	

Labeler - Delta Brands, Inc (102672008)

Revised: 3/2020

Delta Brands, Inc