

**HYGIENE CLEAN EUCALYPTUS HAND SANITIZER- benzalkonium chloride liquid
USA Broom LLC**

Hygiene Clean Eucalyptus Hand Sanitizer

Drug Facts

Active Ingredient

Benzalkonium Chloride 0.13%

Purpose

Antimicrobial

Uses

- For hand sanitizing to decrease the bacteria on skin.
- Recommended for repeated use.

Warnings

For external use only

When using this product avoid

contact with eyes. In case of eye contact, flush eyes with water.

Stop use and ask a doctor if

irritation or redness develops, or if 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

- Pump a small amount of foam into palm of hand
- Rub thoroughly over all surfaces of both hands
- Rub hands together briskly until dry

Inactive ingredients:

Water; Propylene Glycol; Lauramine Oxide; Undeceth-7; Disodium EDTA; Aloe Barbadensis Gel; Glycereth-2-Cocotate; DMDM Hydantoin; Fragrance, Citric Acid

Package Labeling:50ml



HAND SANITIZER
ON-THE-GO & CAR SAFE*
ALCOHOL-FREE
KILLS 99% OF ALL GERMS.
1.7 FL. OZ (50mL)

MADE IN U.S.A.

* Hygiene Clean will not degrade at temperatures up to 212F.

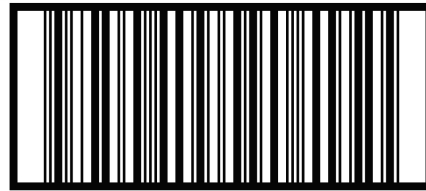


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**HYGIENE CLEAN**
Manufactured for USA Broom, LLC
www.hygieneclean.com
901 S 2nd Avenue Dodge City, KS 67801
620-682-7133
TRA-2OZ-EUC



Package Labeling:50ml 24 bottles



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24 - 1.7 oz.
(50 mL) Bottles
Net Contents - 40.8 oz.
(0.32 gal) (1.2 L)

TRA-2OZ-EUC

HYGIENE CLEAN EUCALYPTUS HAND SANITIZER

benzalkonium chloride liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:80499-013
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII: 7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	1.3 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
LAURAMINE OXIDE (UNII: 4F6FC4MI8W)	
UNDECETH-7 (UNII: R6B5PCO2JN)	
EDETATE DISODIUM ANHYDROUS (UNII: 8NLQ36F6MM)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
GLYCERETH-2 COCOATE (UNII: JWM00VS7HC)	

DMDM HYDANTOIN (UNII: BYR0546TOW)

CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:80499-013-01	50 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/30/2020	
2	NDC:80499-013-02	24 in 1 BOX	10/30/2020	
2		50 mL in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	10/30/2020	

Labeler - USA Broom LLC (117638854)

Revised: 10/2023

USA Broom LLC