HYGIENE CLEAN BUBBLE GUM HAND SANITIZER- benzalkonium chloride liquid USA Broom LLC

Hygiene Clean Bubble Gum 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-Cocoate; DMDM Hydantoin; Fragrance, Citric Acid

Package Labeling:50ml



Package Labeling:50ml 24 bottles





HYGIENE° CLEAN

Manufactured for USA Broom, LLC www.hygieneclean.com 901 S 2nd Avenue Dodge City, KS 67801 620-682-7133 24 - 1.7 oz. (50 mL) Bottles Net Contents - 40.8 oz (0.32 gal) (1.2 L)

TRA-2OZ-BUB

Drug Facts		
Uses - For hand sanitizing to decrease the bacteria on skin Recommended for repeated use.		
Warnings For external use only		

Stop use and ask a doctor if irritation or redness develops, or if condition persists fo

KEEP OUT OF REACH OF CHILDREN

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

DIRECTIONS

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HYGIENE CLEAN BUBBLE GUM HAND SANITIZER

benzalkonium chloride liquid

Product Information Product Type HUMAN OTC DRUG Item Code (Source) NDC:80499-012 Route of Administration TOPICAL

Active Ingredient/Active Moiety Ingredient Name BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y) BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII: 7N6JUD5X6Y) BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII: 7N6JUD5X6Y)

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)	

P	Packaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
1	NDC:80499-012- 01	50 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/30/2020			
2	NDC:80499-012- 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