ENVIROPURE ANTIBACTERIAL HAND SANITIZER MAXIMUM STRENGTH (ALCOHOL-FREE)- benzalkonium chloride liquid
Windmill Health Products, 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.

EnviroPure™ Antibacterial Hand Sanitizer Maximum Strength (Alcohol-Free)

Drug Facts

Active ingredient

Benzalkonium Chloride 0.13%

Purpose

Antimicrobial and First aid antiseptic

Uses

- For hand sanitizing to decrease bacteria on the skin. Recommended for repeated use.
- For wound sanitizing to help prevent bacterial contamination in cuts, scrapes, burns, lacerations and skin infections.

Warnings

For external use only.

When using this product avoid contact with eyes. In case of eye contact, flush eyes with water.

Discontinue use if irritation or redness develops. If condition persists for more than 72 hours consult a doctor.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions

- Pump onto hands as needed. Rub briskly together until dry.
- Pump onto wounds 3 times a day after cleaning. Allow foam to dissipate. Wipe excess with sterile gauze. May be bandaged once dry.

Inactive ingredients

ionized water, urea

Distributed by: Windmill Health Products®
10 Henderson Drive
West Caldwell, NJ 07006

PRINCIPAL DISPLAY PANEL - 1 gal Bottle Label

ENVIROPURE™

ANTIBACTERIAL

HAND
SANITIZER

MAXIMUM STRENGTH

Eliminates 99.99% of Germs

alcohol-free
triclosan-free & fragrance-free

BENZALKONIUM CHLORIDE 0.13%

BAC-D
BACTERIAL DEFENSE®

128 fl. oz. (1 gal)

ENVIROPURE ANTIBACTERIAL HAND SANITIZER MAXIMUM STRENGTH (ALCOHOL-FREE)

benzalkonium chloride liquid

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:74154-004
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 mg in 100 mL

Inactive Ingredients

Ingredient Name	Strength
water (UNII: 059QF0KO0R)	
urea (UNII: 8W8T17847W)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:74154-004-10	3785.41 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/23/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	04/23/2020	

Labeler - Windmill Health Products, LLC (831136267)

Establishment

Name	Address	ID/FEI	Business Operations
Cemi International		015038336	MANUFACTURE(74154-004)

Revised: 2/2022 Windmill Health Products, LLC