BENZALKONIUM CHLORIDE- benzalkonium chloride 0.13% soap
HEB

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.

HEB 279.003/279AD Antibacterial Hand Soap

Active ingredient

Benzalkonium chloride 0.13%

Purpose

Antibacterial

Use

for handwashing to decrease bacteria on the skin

Warnings

For external use only: hand only

When using this product

- avoid contact with eyes. If contact occurs, rinse eyes thoroughly with water

Stop use and ask a doctor if

- irritation or redness develops
- 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
- apply palmful to hands
- scrub thoroughly
- rinse thoroughly

Inactive ingredients

water, cocamidopropyl betaine, lauramine oxide, PEG-150 distearate, sodium chloride, cetrimonium chloride, decyl glucoside, glycerin, fragrance, distearet-75 IPID, citric acid, tetrasodium EDTA, DMDM hydantoin, benzophenone-4, yellow 5, red 40, red 33

Questions?

Call 1-888-593-0593

Distributed by HEB San Antonio, TX 78204

Made in the USA of US and Imported Parts

We hope you are satisfied with this product. If not, we will cheerfully refund your money. Lot number: on package. 1-888-593-0593

principal display panel

Hill Country Essentials

Antibacterial Hand Soap

Moisturizing

7.5 FL OZ (221 mL)

BENZALKONIUM CHLORIDE

benzalkonium chloride 0.13% soap

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:37808-278

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 g in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
LAURAMINE OXIDE (UNII: 4F6FC4MI8W)	
PEG-150 DISTEARATE (UNII: 6F36Q0I0AC)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
CETRIMONIUM CHLORIDE (UNII: UC9PE95IBP)	
DECYL GLUCOSIDE (UNII: Z17H97EA6Y)	
GLYCERIN (UNII: PDC6A3C0OX)	
DISTEARETH-75 ISOPHORONE DIISOCYANATE (UNII: 5365FJ30SC)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
EDETATE SODIUM (UNII: MP1J8420LU)	
DMDM HYDANTOIN (UNII: BYR0546TOW)	
SULISOBENZONE (UNII: 1W6L629B4K)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:37808-278-96	221 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	08/03/2016	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	08/03/2016	

Labeler - HEB (007924756)

Registrant - Vi-Jon, LLC (790752542)

Establishment

Name	Address	ID/FEI	Business Operations
Vi-Jon, LLC		088520668	manufacture(37808-278)

Establishment

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Vi-Jon, LLC		790752542	manufacture(37808-278)

Revised: 12/2022 HEB