

**MOISTURIZING ANTIBACTERIAL- benzalkonium chloride 0.13% soap
Vi-Jon, 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.

Germ-X 842.000/842AA

Inactive ingredient

Benzalkonium chloride 0.13%

Purpose

Antibacterial

Use

for handwashing to decrease bacteria on the skin

Warnings

For external use only: hands 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, lauramine oxide, cocamidopropyl betaine, lauramidopropylamine oxide, sodium chloride, myristamidopropylamine oxide, glycerin, fragrance, disteareth-75 IPDI, PEG-150 distearate, citric acid, tetrasodium EDTA, benzophenone-4, sodium benzoate, Citrus nobilis (mandarin orange) oil, Camellia sinensis leaf extract, Zingiber officinale (ginger) root oil, blue 1, yellow 5

disclaimer

This product is not manufactured or distributed by Colgate-Palmolive Company., distributor of Softsoap Antibacterial Hand Soap with Moisturizers Fresh Citrus

Adverse Reactions

Manufactured by: Vi-Jon, LLC.,

St. Louis, MO 63114

Questions or Comments? 1-888-593-0593

principal display panel

germ-X

SINCE 1997

ANTIBACTERIAL

HAND SOAP

WATERFALL SCENT

WASHES AWAY GERMS

pH Balanced,

Dermatologist Tested

FORMULA MADE IN USA

EMPLOYEE-OWNED

16FL OZ (473 mL)



MOISTURIZING ANTIBACTERIAL

benzalkonium chloride 0.13% soap

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:11344-842
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: 059QF0K00R)	
LAURAMINE OXIDE (UNII: 4F6FC4MI8W)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
LAURAMIDOPROPYLAMINE OXIDE (UNII: I6KX160QTV)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
MYRISTAMIDOPROPYLAMINE OXIDE (UNII: 3HSF539C9T)	
GLYCERIN (UNII: PDC6A3C0OX)	
DISTEARETH-75 ISOPHORONE DIISOCYANATE (UNII: 5365FJ30SC)	
PEG-150 DISTEARATE (UNII: 6F36Q0I0AC)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
EDETATE SODIUM (UNII: MP1J8420LU)	
SULISOBENZONE (UNII: 1W6L629B4K)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
MANDARIN OIL (UNII: NJO720F72R)	
GREEN TEA LEAF (UNII: W2ZU1RY8B0)	
GINGER OIL (UNII: SAS9Z1SVUK)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11344-842-43	473 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	04/05/2021	
2	NDC:11344-842-32	354 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	04/05/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	04/05/2021	

Labeler - Vi-Jon, LLC (790752542)

Registrant - Vi-Jon, LLC (790752542)

Establishment

Name	Address	ID/FEI	Business Operations
Vi-Jon, LLC		790752542	manufacture(11344-842)

Establishment

Name	Address	ID/FEI	Business Operations
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Vi-Jon, LLC

088520668

manufacture(11344-842)

Revised: 4/2022

Vi-Jon, LLC