HAND WASH- benzalkonium chloride liquid United Natural Foods, Inc. dba UNFI

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.

Antibacterial Foaming Hand Wash 466.001/466AB

Active 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 dry hands
- scrub thoroughly
- rinse thoroughly

Inactive ingredients

water, cocamidopropyl betaine, lauramidopropylamine oxide, lauramine oxide, myristamidopropylamine oxide, glycerin, fragrance, citric acid, tetrasodium EDTA, benzophenone-4, sodium benzoate, red 4, yellow 5

Adverse reactions

DISTRIBUTED BY SUPERVALU INC., EDEN PRAIRIE, MN 55344 USA 877-932-7948 www.supervalu-ourownbrands.com

Principal display panel

EQUALINE

ANTIBACTERIAL

Foaming

Hand Wash

Original

7.5 FL OZ (221 mL)



HAND WASH

benzalkonium chloride liquid

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:41163-941	
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)			
COCAMIDOPROPYL BETAINE (UNII: 50CF3011KX)			
LAURAMIDOPROPYLAMINE OXIDE (UNII: I6KX160QTV)			
LAURAMINE OXIDE (UNII: 4F6FC4MI8W)			
MYRISTAMIDOPROPYLAMINE OXIDE (UNII: 3HSF539C9T)			
GLYCERIN (UNII: PDC6A3C0OX)			
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)			
EDETATE SODIUM (UNII: MP1J8420LU)			
SULISOBENZONE (UNII: 1W6L629B4K)			
FD&C RED NO. 4 (UNII: X3W0AM1JLX)			
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)			

l	Packaging				
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
		NDC:41163- 941-96	221 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/15/2016	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	03/15/2016	

Labeler - United Natural Foods, Inc. dba UNFI (943556183)

Registrant - Vi-Jon, LLC (790752542)

Establishment			
Name	Address	ID/FEI	Business Operations
Vi-Jon, LLC		088520668	manufacture(41163-941)

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Name	Address	ID/FEI	Business Operations
Vi-Jon, LLC		790752542	manufacture(41163-941)

Revised: 5/2023 United Natural Foods, Inc. dba UNFI