INOFOAM ANTIBACTERIAL FOAMING HAND WASH- benzalkonium chloride liquid
Avro Enterprises, LLC

Inofoam Antibacterial Foaming Hand Wash

Drug Facts

Active ingredient

Benzalkonium Chloride 0.13%

Purppose

Antiseptic

Uses

- Handwash to help reduce bacteria on the skin that potentially can cause disease.
- Recommended for repeated use.

Warings

For external use only.

When using this product

Keep out of eyes. In case of eye contact, flush eyes with water.

Stop use and ask a doctor if

irritation and redness develop 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.

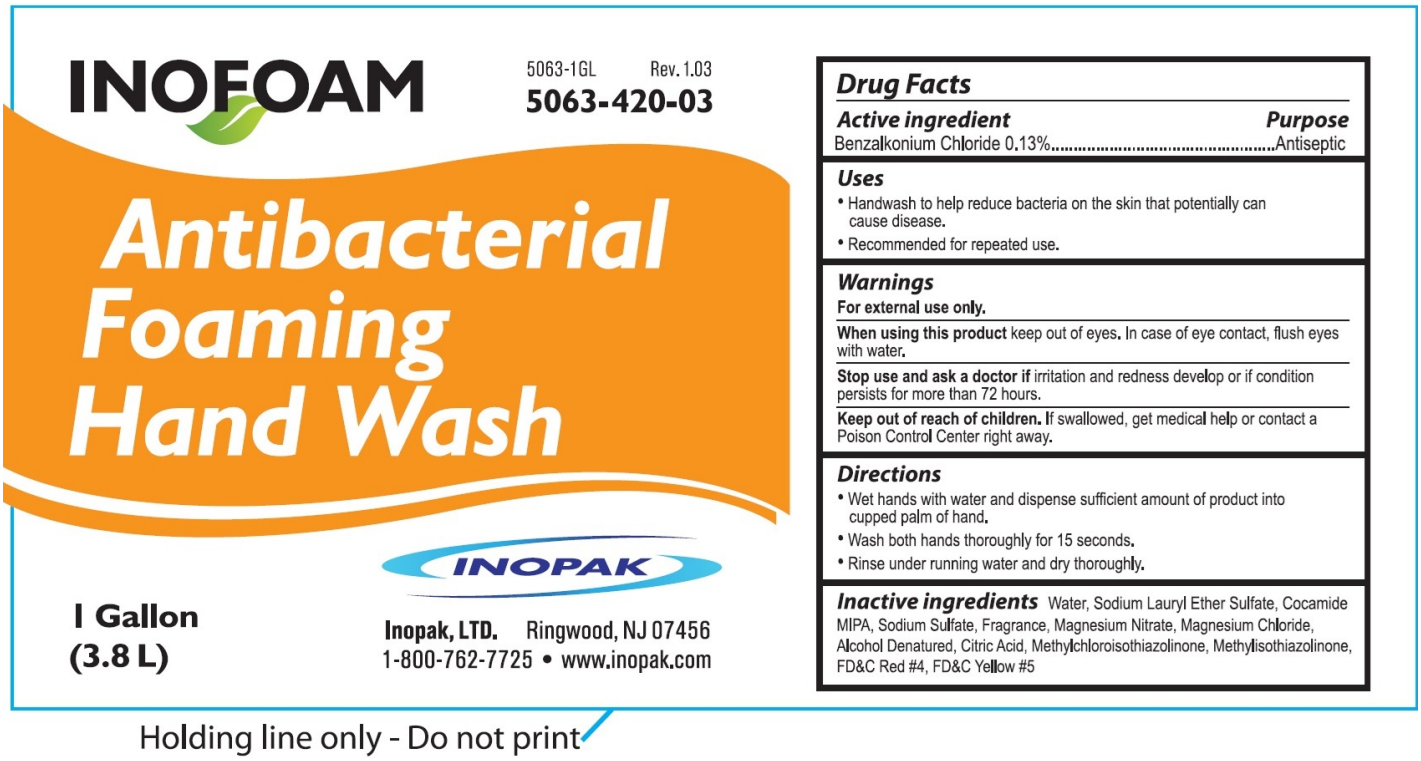
Directions

• Wet hands with water and dispense sufficient amount of product into cupped palm of hand. • Wash both hands thoroughly for 15 seconds. • Rinse under running water and dry thoroughly.

Inactive ingredients

Water, Sodium Lauryl Ether Sulfate, Cocamide MIPA, Sodium Sulfate, Fragrance, Magnesium Nitrate, Magnesium Chloride, Alcohol Denatured, Citric Acid, Methylchloroisothiazolinone, Methylisothiazolinone, FD&C Red#4, FD&C Yellow #5

Package Labeling:



INOFOAM ANTIBACTERIAL FOAMING HAND WASH

benzalkonium chloride liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:73062-032
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)	
SODIUM LAURETH-3 SULFATE (UNII: BPV390UAP0)	
COCO MONOISOPROPANOLAMIDE (UNII: 21X4Y0VTB1)	
SODIUM SULFATE (UNII: 0YPR65R21J)	
MAGNESIUM NITRATE (UNII: 77CBG3UN78)	
MAGNESIUM CHLORIDE (UNII: 02F3473H9O)	
ALCOHOL (UNII: 3K9958V90M)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	

METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
FD&C RED NO. 4 (UNII: X3W0AM1JLX)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73062-032-01	1 in 1 CARTON	11/20/2023	
1		3800 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)	11/20/2023	

Labeler - Avro Enterprises, LLC (804030166)

Establishment

Name	Address	ID/FEI	Business Operations
Fuller Industries, Inc		078704329	manufacture(73062-032)

Revised: 11/2023 Avro Enterprises, LLC