

**CHEMSTAR INSTANTFOAM NONALCOHOL SANITIZER- benzalkonium chloride liquid
Deb USA, Inc.**

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.

Active ingredient

Benzalkonium Chloride, 0.13%

Purpose

Antibacterial

Uses

For hand sanitizing to reduce bacteria on the skin

Warnings

For external use only

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

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center right away.

Directions

Apply one shot to dry hands, rub into skin

No rinsing required

Inactive ingredients

Water, Propylene Glycol, Aloe Barbadensis Leaf Juice, Cocamidopropyl Betaine, Lauramine Oxide, Tetrasodium EDTA, Fragrance, Citric Acid, Magnesium Nitrate, Methylchloroisoithiazolinone, Magnesium Chloride, Methylisothiazolinone, Blue 1 (CI 42090), Red 33 (CI 17200)

Chemstar Non-Alcohol Foaming Hand Sanitizer

CHEMSTAR

Non-Alcohol Foaming Hand Sanitizer

CHEMSTAR

Integrated Food Safety Solutions

120 Interstate West Parkway

Suite 100

Lithia Springs, GA 30122

www.chemstarcorp.com

1L

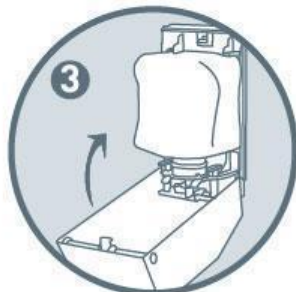
33.8 Fl. Oz.

Non-Alcohol Foaming Hand Sanitizer

Rev. 12-12

CHEMSTAR

Non-Alcohol Foaming Hand Sanitizer



Drug Facts

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33.8 Fl. Oz.

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Rev. 12-12

CHEMSTAR INSTANTFOAM NONALCOHOL SANITIZER

benzalkonium chloride liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:11084-111
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)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
LAURAMINE OXIDE (UNII: 4F6FC4MI8W)	
EDETATE SODIUM (UNII: MP1J8420LU)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
MAGNESIUM NITRATE (UNII: 77CBG3UN78)	
MAGNESIUM CHLORIDE (UNII: 02F3473H9O)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11084-111-27	1000 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	12/01/2012	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	12/01/2012	

Labeler - Deb USA, Inc. (607378015)**Registrant** - Deb USA, Inc. (607378015)**Establishment**

Name	Address	ID/FEI	Business Operations
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Deb USA, Inc.

078805627

manufacture(11084-111)

Revised: 10/2017

Deb USA, Inc.