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, Methylchloroisothiazolinone, Magnesium Chloride, Methylisothiazolinone, Blue 1 (CI 42090), Red 33 (CI 17200)

Chemstar Non-Alcohol Foaming Hand Sanitizer

Chemstar

CHEMSTAR

Non-Alcohol Foaming **Hand Sanitizer**



Drug Facts

Active ingredient

Purpose

Benzalkonium Chloride, 0.13% Antibacterial

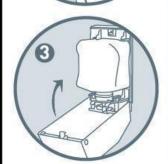
Uses For hand sanitizing to reduce bacteria on the skin

Warnings

For external use only



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, Methylchloroisothiazolinone, Magnesium Chloride, Methylisothiazolinone, Blue 1 (Cl 42090), Red 33 (Cl 17200)



120 Interstate West Parkway Suite 100 Lithia Springs, GA 30122 1-800-327-0777 www.chemstarcorp.com

33.8 Fl. Oz.

Non-Alcohol Foaming Hand Sanitizer



CHEMSTAR INSTANTFOAM NONALCOHOL SANITIZER

| Product Information | | | | |
|-------------------------|----------------------|--------------------|---------------|--|
| Product Type | HUMAN OTC DRUG LABEL | Item Code (Source) | NDC:11084-111 | |
| Route of Administration | TOPICAL | DEA Sche dule | | |

| | Active Ingredient/Active Moiety | | | |
|---|--------------------------------------|-----------------------|----------------|--|
| ı | Ingredient Name | Basis of Strength | Strength | |
| ı | BENZALKONIUM CHLORIDE (BENZALKONIUM) | BENZALKONIUM CHLORIDE | 1.3 mg in 1 mL | |

| Inactive Ingredients | | |
|-----------------------------|----------|--|
| Ingredient Name | Strength | |
| WATER | | |
| PROPYLENE GLYCOL | | |
| ALOE VERA LEAF | | |
| COCAMIDO PRO PYL BETAINE | | |
| LAURAMINE O XIDE | | |
| EDETATE SO DIUM | | |
| ANHYDRO US CITRIC ACID | | |
| D&C RED NO. 33 | | |
| FD&C BLUE NO. 1 | | |
| METHYLCHLOROISOTHIAZOLINONE | | |
| METHYLISO THIAZO LINO NE | | |
| MAGNESIUM NITRATE | | |
| MAGNESIUM CHLO RIDE | | |

| P | ackaging | | | |
|---|------------------|------------------------------|----------------------|--------------------|
| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
| 1 | NDC:11084-111-27 | 1000 mL in 1 BOTTLE, PLASTIC | | |

| Marketing Information | | | | |
|-------------------------|--|----------------------|--------------------|--|
| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date | |
| OTC monograph not final | part333A | 12/0 1/20 12 | | |
| | | | | |

Labeler - Deb USA, Inc. (607378015)

Registrant - Deb USA, Inc. (607378015)

| Establishment | | | | | |
|---------------|---------|-----------|------------------------|--|--|
| Name | Address | ID/FEI | Business Operations | | |
| Deb USA, Inc. | | 607378015 | manufacture(11084-111) | | |

Revised: 12/2012 Deb USA, Inc.