DEFENDER ANTISEPTIC FOAM HAND SANITIZER ALCOHOL-FREE- benzalkonium chloride liquid
Scientific Molecular Technologies

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.

DEFENDER Antiseptic Foam Hand Sanitizer Alcohol-Free

Drug Facts

Active Ingredients

Benzalkonium Chloride 0.13%

Purpose

Antimicrobial

Uses

- For hand sanitizing to decrease bacteria on the skin.
- Recommended for repeated use.

Warnings

- **For external use only.**

Do not use in eyes

if contact occurs, flush eyes with water.

Stop use and ask a doctor

if irritation or redness develops. If condition persists for more than 72 hours, consult a doctor.

Keep out of reach of children

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

Directions

- Pump a small amount of foam into palm of hand.
- Wet hands thoroughly with product and allow to dry without wiping.
- Rub hands together briskly until dry.

Inactive Ingredients

Water, dihydroxypropyl PEG-5 linoleammonium chloride, glycereth-2 cocoate, behentrimonium chloride, dihydroxyethyl cocamine oxide, fragrance

DE Fender SCIENTIFIC MOLECULAR TECHNOLOGIES ANTISEPTIC FOAM Hand Sanitizer ALCOHOL-FREE ELIMINATES 99.999% OF MOST COMMON GERMS THAT CAUSES ILLNESS NO RINSE MOISTURIZES LEAVES SKIN SOFT EFFECTIVE AGAINST STAPH, MRSA, VRE, NOROVIRUS 1.7 fl oz (50 ml)

Product Label

DEFENDER ANTISEPTIC FOAM HAND SANITIZER ALCOHOL-FREE

benzalkonium chloride liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:42953-002
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)	
DIHYDROXYPROPYL PEG-5 LINOLEAMMONIUM CHLORIDE (UNII: 0Y0NQR2GH1)	
GLYCERETH-2 COCOATE (UNII: JWM00VS7HC)	
BEHENTRIMONIUM CHLORIDE (UNII: X7GNG3S47T)	
DIHYDROXYETHYL COCAMINE OXIDE (UNII: 8AR51R3BL5)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:42953-002-00	50 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	10/11/2012	
2	NDC:42953-002-01	550 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	10/11/2012	
3	NDC:42953-002-02	1000 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	10/11/2012	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	10/11/2012	

Labeler - Scientific Molecular Technologies (028159474)

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