ANTIBACTERIAL HAND MOUSSE - benzalkonium chloride liquid MAYBORN 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.

Drug Facts

Active ingredient

Benzalkonium Chloride 0.13%

Purpose

Antimicrobial

Uses

For hand washing to decrease bacteria on skin.

Recommended for repeated use.

Warnings

For external use only.

When using this product avoid contact with eyes.

In case of eye contact, flush eyes with water.

Stop use and ask a doctor if irritation or redness develops, 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

Pump a small amount of foam into the palm of hand

Rub thoroughly over all surfaces of both hands for 15 seconds

Rinse with potable water.

Inactive ingredients

Water, coco-glucoside, laurtrimonium chloride, cocamidopropylamine oxide.

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closer to nature

ANTIBACTERIAL

HAND SOAP

MOUSSE

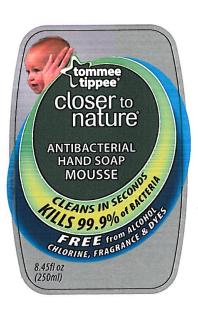
CLEANS IN SECONDS

KILLS 99.9% of BACTERIA

FREE from ALCOHOL

CHLORINE, FRAGRANCE and DYES

8.45fl oz



ANTIBACTERIAL HAND MOUSSE

benzalkonium chloride liquid

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Product	Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:76273-002

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKO NIUM CHLO RIDE (UNII: F5UM2KM3W7) (BENZALKO NIUM -	BENZALKONIUM	0.13 mL
UNII:7N6JUD5X6Y)	CHLORIDE	in 100 mL

Inactive Ingredients

inactive ingredients		
Ingredient Name	Strength	
WATER (UNII: 059QF0KO0R)		
COCO GLUCOSIDE (UNII: ICS790225B)		
COCAMIDO PRO PYLAMINE O XIDE (UNII: M4SL8217HK)		

Packaging

П	- ucing			
l	# Item Code	Package Description	Marketing Start Date	Marketing End Date
l	1 NDC:76273-002-03	250 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	07/28/2011	

Labeler - MAYBORN USA, INC. (031274363)

Establishment				
Name	Address	ID/FEI	Business Operations	
ORGANICA (UK) LTD		235330151	manufacture	

Revised: 8/2011 MAYBORN USA, INC.