NATIVE GREEN FOAMING HAND SANITIZER- benzalkonium chloride solution
Native Green

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 0.1%

Purpose

Antimicrobial

Uses

- For hand sanitizing to decrease bacteria on the skin

Warning

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 palm of hand
- Rub thoroughly over all surfaces of both hands
- Rub hands together briskly until dry

Inactive Ingredients Water, dihydroxpropyl PEG-5 linoleammonium chloride, glycereth-2 cocoate, behentrimonium chloride, dihydroxyethyl cocamine oxide, fragrance

Principal Display Panel – Bottle Label

NATIVE GREEN
FOAMING HAND SANITIZER
ALCOHOL-FREE – WITH MOISTURIZERS

Cleaner, Greener

SEE SIDE PANEL FOR ADDITIONAL INFORMATION.

NATIVE GREEN FOAMING HAND SANITIZER

benzalkonium chloride solution

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:50409-514
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
benzalkonium chloride (UNII: F5UM2KM3W7) (benzalkonium - UNII:7N6JUD5X6Y)	benzalkonium chloride	1 g in 1000 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:50409-514-82	550 mL in 1 BOTTLE		
2	NDC:50409-514-42	50 mL in 1 BOTTLE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333	04/21/2010	

Labeler - Native Green (791610038)

Establishment

Name	Address	ID/FEI	Business Operations
Canberra Corporation		068080621	MANUFACTURE

Revised: 4/2010 Native Green