PURACY NATURAL FOAMING HAND SANITIZER- benzalkonium chloride solution Puracy LLC

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

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

Inactive ingredients

Water, cetrimonium chloride, laurtrimonium chloride, dihydroxyethyl cocamine oxide, glycereth-17 cocoate, citric acid

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Leaves skin feeling silky soft Fast-drying, non-sticky, mess-free foam
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PURACY NATURAL FOAMING HAND SANITIZER						
benzalkonium chloride solution						
Product Information						
Product Type	HUMAN OTC DRUG	Item Code (Sou	ırce)	NDC:7290	5-001	
Route of Administration	TOPICAL					
Active Ingredient/Active Moiety						
Ingredient Name Basis of Strength Strength					Strength	

BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM -	BENZALKONIUM	0.1 g
UNII:7N6JUD5X6Y)	CHLORIDE	in 100 mL

Inactive Ingredients	
Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
CETRIMONIUM CHLORIDE (UNII: UC9PE95IBP)	
LAURTRIMONIUM CHLORIDE (UNII: A8 1MS 10 FIC)	
DIHYDROXYETHYL COCAMINE OXIDE (UNII: 8AR51R3BL5)	
GLYCERETH-17 COCOATE (UNII: 3057VPT0KC)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72905-001- 08	251 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	0 2/20 /20 19	
2	NDC:72905-001- 05	18927 mL in 1 PAIL; Type 0: Not a Combination Product	0 2/20 /20 19	
Marketing Information				

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	02/20/2019	

Labeler - Puracy LLC (079535215)

Establishment

Name	Address	ID/FEI	Business Operations
Bayscience Formulators, LLC		162930544	manufacture(72905-001)

Revised: 2/2019

Puracy LLC