

**APOTHECARY HOME- benzalkonium chloride liquid
HOME & BODY COMPANY**

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.

APOTHECARY HOME SOAP

Active ingredient

Benzalkonium Chloride 0.13%

Purpose

Antimicrobial

Use

help eliminate bacteria on hands

Warnings

For external use only.

When using this product do not use in or near the eyes.

In case of contact, rinse eyes thoroughly with water.

Keep out of reach of children, except under adult supervision. If swallowed, get medical help or contact a Poison Control Center right away.

Directions

Wash hands and rinse

Inactive ingredients

Water (Aqua), Sodium Laureth Sulfate, Cocamidopropyl Betaine, Sodium Chloride, Sodium Benzoate, Phenoxyethanol, Polysorbate 20, Fragrance, Glycerin

APOTHECARY HOME	ANTIBACTERIAL HAND SOAP GENTLY CLEANS AND PROTECTS	APOTHECARY HOME	Drug Facts
	FORMULATED AND DESIGNED BY: HOB Co.  		Active Ingredient Purpose 0.13% Benzalkonium chlorideAntibacterial
HOME & BODY COMPANY HUNTINGTON BEACH, CA 92648 WWW.HOMEANDBODYCO.COM	 1 80969 00082 9 This area is reserved for the expiration date and batch number	With the power of 0.13 percent Benzalkonium Chloride, this hand soap reduces the bacteria on the skin, gently cleans and moisturizes, and leaves the skin smelling great. KILLS 99.99% OF GERMS	Use helps eliminate bacteria on hands
			BATCH NO. 01
ANTIBACTERIAL HAND SOAP			Directions wash hands and rinse
LAVENDER SAGE			Inactive Ingredients Water (Aqua), Sodium Laureth Sulfate, Cocamidopropyl Betaine, Sodium Chloride, Sodium Benzoate, Phenoxyethanol, Polysorbate 20, Fragrance, Glycerin
e 636ml HOME & BODY COMPANY 21.5fl.oz.		<small>Home and Body Company 5900 SkyLab Ste. 100, Huntington Beach, WWW.HOMEANDBODYCO.COM</small>  	

APOTHECARY HOME

benzalkonium chloride liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:73746-005
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)	
SODIUM LAURETH SULFATE (UNII: BPV390UAP0)	
CO CAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
GLYCERIN (UNII: PDC6A3C0OX)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73746-005-01	636 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/23/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	03/23/2020	

Labeler - HOME & BODY COMPANY (081290720)

Establishment

Name	Address	ID/FEI	Business Operations
HOME & BODY COMPANY		081290720	manufacture(73746-005)

Revised: 3/2020

HOME & BODY COMPANY