

ACTIVE GUARD- 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.

Active Guard Antibacterial Hand Soap 21.5 fl.oz

Active ingredient

0.13% Benzalkonium Chloride

Purpose

Antibacterial

Use

Helps reduce bacterial 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

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), Cocamidopropyl Betaine, Lauramine Oxide, Cellulose Gum, DMDM Hydantoin, Polysorbate 20, Fragrance, Glycerin

Product label



ANTIBACTERIAL HAND SOAP

KILLS 99.9% OF GERMS

**0.13%
Benzalkonium Chloride**

- **Antibacterial**
- **Moisturizing**
- **Gentle Formula**

Net Wt. 636 ml / 21.5 fl.oz

Drug Facts

Active Ingredient	Purpose
0.13% Benzalkonium Chloride.....	Antibacterial

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Directions

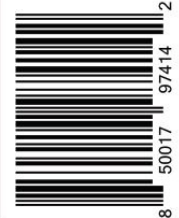
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Inactive Ingredients

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Home and Body Company

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United States (USA)
www.homeandbodyco.com #06AGHS



EXP. XX/2021
This area is reserved for
batch number
NDC # 73746-018-01

ACTIVE GUARD

benzalkonium chloride liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:73746-018
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
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
CARBOXYMETHYLCELLULOSE SODIUM (UNII: K679OBS311)	
LAURAMINE OXIDE (UNII: 4F6FC4MI8W)	
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
DMDM HYDANTOIN (UNII: BYR0546TOW)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73746-018-01	636 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/05/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	05/05/2020	

Labeler - HOME & BODY COMPANY (081290720)

Registrant - HOME & BODY COMPANY (081290720)

Establishment

Name	Address	ID/FEI	Business Operations
HOME & BODY COMPANY		081290720	manufacture(73746-018)

Revised: 5/2020

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