

**ACTIVE GUARD REFILL- 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.*

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**Active Guard Antibacterial Hand Soap Refill 64 fl.oz**

**Active ingredient**

0.13% Benzalkonium Chloride

**Purpose**

Antibacterial

**Use**

Helps reduce bacterial on hands

**Warning**

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 REFILL

**KILLS 99.9% OF GERMS**

**0.13%**  
**Benzalkonium Chloride**

## Drug Facts

Active Ingredient	Purpose
0.13% Benzalkonium chloride	Antibacterial

**Use** Helps reduce bacteria on hands

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### Home and Body Company

5900 SKYLAB RD, HUNTINGTON BEACH, California (CA)  
92647, United States (USA). WWW.HOMEANDBODYCO.COM

**Net Wt. 1893 ml / 64 fl.oz**

EXP. 11/2021  
BATCH. 20135370AGARHS

NDC # 73746-017-01



## ACTIVE GUARD REFILL

benzalkonium chloride liquid

### Product Information

Product Type

HUMAN OTC DRUG

Item Code (Source)

NDC:73746-017

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)	
DMDM HYDANTOIN (UNII: BYR0546TOW)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
LAURAMINE OXIDE (UNII: 4F6FC4M18W)	
WATER (UNII: 059QF0K00R)	
CARBOXYMETHYLCELLULOSE SODIUM (UNII: K679OBS311)	
GLYCERIN (UNII: PDC6A3C0OX)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73746-017-01	1893 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(73746-017)

Revised: 5/2020

HOME & BODY COMPANY