HAND RX ANTIBACTERIAL HAND- benzalkonium chloride liquid
BLUE CROSS LABORATORIES, INC.

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 0.13 percent

Purpose

Antibacterial

Uses:

For hand washing to decrease bacteria on the skin.

Warnings:

For external use only.

When using this product Avoid contact with eyes. In case of eye contact, flush with water.

Stop use and ask a doctor if

irritation and redness develops.

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center right away.

Directions

- Wet hands as needed
- Lather vigorously for at least 15 seconds
- Wash skin, rinse and dry thoroughly.

Inactive ingredients

Water (Aqua), Cocamidopropyl Betaine, Sodium Cocoamphoacetate, PEG-150 Distearate, Fragrance, PEG-40 Hydrogenated Castor Oil, Citric Acid, Methylisothiazolinone, Iodopropynyl Butylcarbamate, FD&C Yellow No. 5, FD&C Red No.40, D&C Red No. 33

HAND RX ANTIBACTERIAL HAND

benzalkonium chloride liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:22431-021
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM -	BENZALKONIUM	0.13 g

UNII:7N6JUD5X6Y)	CHLORIDE	in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
SODIUM COCOAMPHOACETATE (UNII: W7Q5E87674)	
IODOPROPYNYL BUTYLCARBAMATE (UNII: 603P14DHEB)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
PEG-150 DISTEARATE (UNII: 6F36Q0I0AC)	
HYDROGENATED CASTOR OIL (UNII: ZF94AP8MEY)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:22431-021-01	355 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/01/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	08/01/2020	

Labeler - BLUE CROSS LABORATORIES, INC. (008298879)

Registrant - BLUE CROSS LABORATORIES, INC. (008298879)

Revised: 8/2020 BLUE CROSS LABORATORIES, INC.