

CLEANHANDS- benzalkonium chloride soap
Hangzhou Huiji Biotechnology Co., Ltd.

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.

73788-103 16.9oz Antibacterial Liquid Hand Soap Unscented (Benzalkonium chloride 0.13%)

Active Ingredient

Benzalkonium Chloride 0.13%

Purpose

Antibacterial

Use

For hand sanitizing to reduce 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 or redness develops.

keep out of reach of children

Keep out of reach of children. If swallowed, get medical help or contact a poison control center right away.

Directions

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

Inactive ingredients

water, cetrimonium chloride, glycerin, lauramidopropylamine oxide, cocamide MEA, sodium chloride, PEG-120 methyl glucose dioleate, citric acid, tetrasodium EDTA, methylchloroisothiazolinone,

methylisothiazolinone.



CLEANHANDS

benzalkonium chloride soap

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:73788-103
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII: 7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.13 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
PEG-120 METHYL GLUCOSE DIOLEATE (UNII: YM0K64F20V)	

EDETATE SODIUM (UNII: MP1J8420LU)	
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
CETRIMONIUM CHLORIDE (UNII: UC9PE95IBP)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
LAURAMIDOPROPYLAMINE OXIDE (UNII: I6KX160QTV)	
COCO MONOETHANOLAMIDE (UNII: C80684146D)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73788-103-01	20 in 1 CARTON	07/15/2020	
1		500 g in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	07/15/2020	

Labeler - Hangzhou Huiji Biotechnology Co., Ltd. (526893497)

Establishment

Name	Address	ID/FEI	Business Operations
Hangzhou Huiji Biotechnology Co., Ltd.		526893497	manufacture(73788-103)

Revised: 2/2022

Hangzhou Huiji Biotechnology Co., Ltd.