

XTRACARE FOAM ANTIBACTERIAL HAND WASH WILD BERRY BLAST- benzalkonium chloride soap

China Ningbo Shangge Cosmetic Technology Corp

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.

XtraCare Foam Antibacterial Hand Wash

Active Ingredient	Purpose
Benzalkonium Chloride 0.1%	Antibacterial

Uses

for hand washing to decrease bacteria on the skin.

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

XtraCare® Foam Antibacterial Hand Wash

Kills 99.9% of bacteria

Compare to Lysol® foam antibacterial hand wash

7.5 Fl Oz (221 mL)

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.

Directions

- pump into hands
- lather vigorously for at least 15 seconds
- rinse hands with water

Inactive Ingredients

water, glycerin, lauramine oxide, PEG-150 distearate, cetrimonium chloride, propylene glycol, fragrance, citric acid, tetrasodium EDTA, cocamide MEA, methylchloroisothiazolinone, methylisothiazolinone, FD&C blue no.1

Other Information: store at room temperature

Questions/Comments? 1-855-345-5575

*This product is not manufactured or distributed by Reckitt Benckiser LLC, the distributor of Lysol® foam antibacterial hand wash.

DISTRIBUTED BY:

REJOICE INTERNATIONAL INC

NORTHVILLE, MI 48168 USA

MADE IN CHINA



XTRACARE FOAM ANTIBACTERIAL HAND WASH WILD BERRY BLAST

benzalkonium chloride soap

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:58503-048
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.2 g in 221 g

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
LAURAMINE OXIDE (UNII: 4F6FC4M18W)	
PEG-150 DISTEARATE (UNII: 6F36Q0I0AC)	
CETRIMONIUM CHLORIDE (UNII: UC9PE95IBP)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	

EDETATE SODIUM (UNII: MP1J8420LU)	
COCO MONOETHANOLAMIDE (UNII: C80684146D)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
FD&C BLUE NO. 1 (UNII: HBR47K3TBD)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:58503-048-01	221 g in 1 BOTTLE; Type 0: Not a Combination Product	01/29/2014	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	01/29/2014	

Labeler - China Ningbo Shangge Cosmetic Technology Corp (529287434)

Establishment

Name	Address	ID/FEI	Business Operations
China Ningbo Shangge Cosmetic Technology Corp		529287434	manufacture(58503-048)

Revised: 9/2019

China Ningbo Shangge Cosmetic Technology Corp