

CARE TOUCH ANTIBACTERIAL HANDSOAP ROSE LAVENDER- benzalkonium chloride liquid

Future Diagnostics Llc

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.

CareTouch ANTIBACTERIAL HAND SOAP ROSE LAVENDER

Drug Facts

Active ingredient

Benzalkonium chloride 0.1

Purpose.....Antibacterial

Uses

For hand washing to decrease bacteria on the skin.

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For hand washing to decrease bacteria on the skin.

Warnings For external use only.

Do not use

- in the eyes.
- if you are allergic to any of the ingredients.

When using this product if eye contact occurs, rinse thoroughly with water.

Stop use and ask a doctor if irritation or redness develops and persists.

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 ingredient

Aqua, Cocamidopropylamine Oxide, Glycerin, Cocamide Methyl MEA, Aloe Barbadensis Leaf Water, PEG-150 Distearate,

Cetrimonium Chloride, Sodium Chloride, Phenoxyethanol, Disodium EDTA, Tocopheryl Acetate.

KILLS 99.99% OF BACTERIA

***kills 99.99% of most germs**

Made in China for Future Brands

Brooklyn, NY 11220 1800.758.3830

caretouchusa.com

Packaging

CareTouch®

ANTIBACTERIAL
HAND SOAP



KILLS 99.9%
OF BACTERIA*

ROSE
LAVENDER

400 mL | 13.52 fl oz

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ITEM# CTHSLR13 caretouchusa.com

CARE TOUCH ANTIBACTERIAL HANDSOAP ROSE LAVENDER

benzalkonium chloride liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:70393-052
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.1 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
CO CAMIDO PROPYLAMINE OXIDE (UNII: M4SL82J7HK)	
GLYCERIN (UNII: PDC6A3C0OX)	
COCOYL METHYL MONOETHANOLAMINE (UNII: 79G1T427CF)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
PEG-150 DISTEARATE (UNII: 6F36Q0I0AC)	
CETRIMONIUM CHLORIDE (UNII: UC9PE95IBP)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
EDETATE DISODIUM ANHYDROUS (UNII: 8NLQ36F6MM)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70393-052-01	400 mL in 1 BOTTLE; Type 0: Not a Combination Product	09/01/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	09/01/2020	

Labeler - Future Diagnostics Llc (080113296)

Establishment

Name	Address	ID/FEI	Business Operations
Shenzhen Lantern Science Co., Ltd.		421222423	manufacture(70393-052)