

ANTIBACTERIAL HANDWASH TROPICAL BEACH- benzalkonium chloride liquid
Ningbo Liyuan Daily Chemical Products 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.

Active Ingredient

Benzalkonium Chloride 0.13%

Purpose

Antibacterial

USES

for hand washing to decrease bacteria on the skin.

WARNING

For external use only.

When using this product avoid contact with eyes. In case of eye contact, flush with clean 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 immediately.

DIRECTIONS

- pump into hands
- lather vigorously for at least 5 seconds
- rinse and dry thoroughly.

Inactive Ingredients:

Water(Aqua),Sodium Laureth Sulfate,Cocamidopropyl Betaine,Coconut Monoethanol Amide,Sodium Chloride,Fragrance,Citric Acid,Methylchloroisothiazolinone,Methylisothiazolinone,FD&C Blue No.1.



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Item# 82742



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Made in China

Distributed By:
Universal Distribution Center
96 Distribution Boulevard,
Edison, NJ 08817

ANTIBACTERIAL HANDWASH TROPICAL BEACH

benzalkonium chloride liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:76176-043
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 mL

Inactive Ingredients

Ingredient Name	Strength
METHYLCHLOROISO THIAZOLINONE (UNII: DEL7T5QRPN)	
METHYLISO THIAZOLINO NE (UNII: 229D0E1QFA)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	

SODIUM CHLORIDE (UNII: 451W47IQ8X)

WATER (UNII: 059QF0KO0R)

SODIUM LAURETH SULFATE (UNII: BPV390UAP0)

COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)

COCO DIETHANOLAMIDE (UNII: 92005F972D)

FD&C BLUE NO. 1 (UNII: HBR47K3TBD)

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:76176-043-01	400 mL in 1 BOTTLE; Type 0: Not a Combination Product	08/01/2018	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	08/01/2018	

Labeler - Ningbo Liyuan Daily Chemical Products Co.,Ltd. (530766098)

Registrant - Ningbo Liyuan Daily Chemical Products Co.,Ltd. (530766098)

Establishment

Name	Address	ID/FEI	Business Operations
Ningbo Liyuan Daily Chemical Products Co.,Ltd.		530766098	manufacture(76176-043)

Revised: 8/2018

Ningbo Liyuan Daily Chemical Products Co.,Ltd.