ANTIBACTERIAL HANDWASH BERRY MEDLEY- 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.

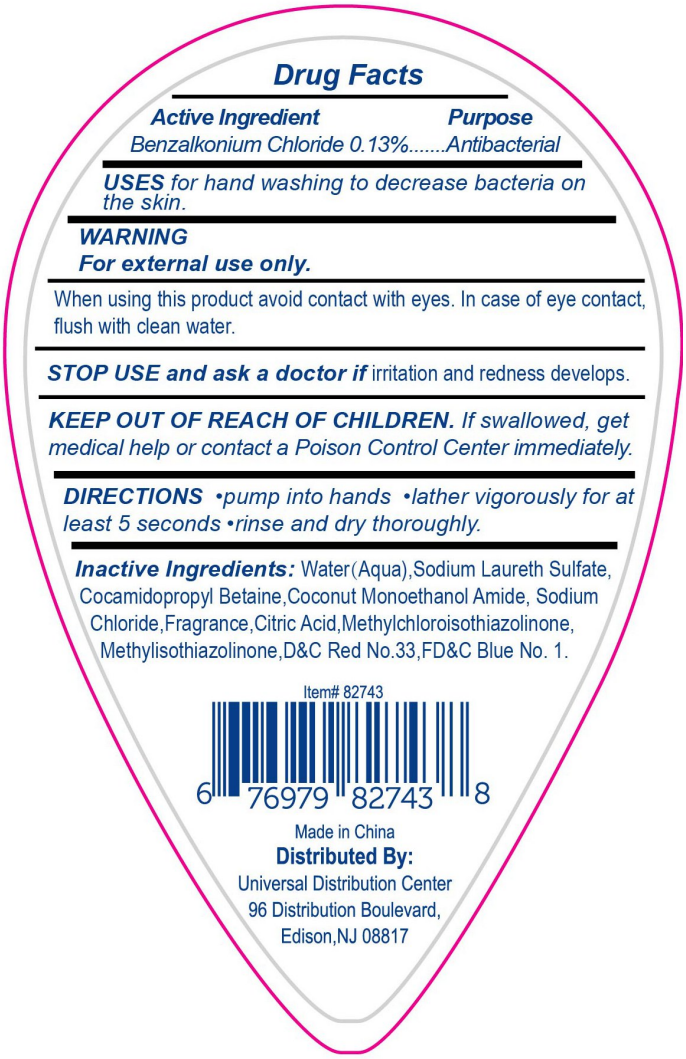
If swallowed, get medical help or contact a Poison Control Center immediately.

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,D&C Red No.33,FD&C Blue No.1.



ANTIBACTERIAL HANDWASH BERRY MEDLEY

benzalkonium chloride liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:76176-044
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**
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
METHYLISOTHIAZOLINONE (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)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:76176-044-01	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(76176-044)

Revised: 8/2018 Ningbo Liyuan Daily Chemical Products Co., Ltd.