

GLOBAL CARE ANTIBACTERIAL HAND ALOE VERA- 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.

Drug Facts

Active ingredient

Benzalkonium Chloride 0.13 percent

Purpose

Antibacterial

Uses

for hand washing to decrease 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 and redness develops.

Keep out of reach of children.

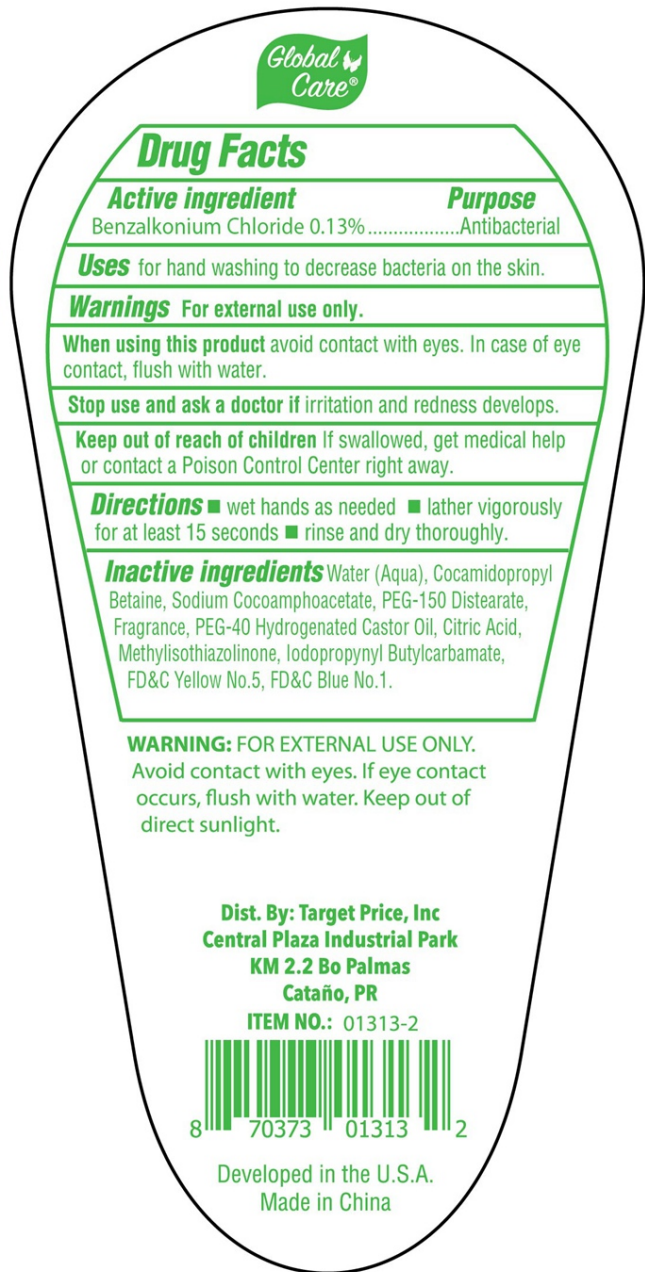
If swallowed, get medical help or contact a Poison Control Center right away.

Directions

- wet hands as needed
- lather vigorously for at least 15 seconds
- rinse and dry thoroughly.

Inactive ingredients

Water (Aqua), Cocamidopropyl Betaine, Sodium Cocoamphoacetate, PEG-150 Distearate, Fragrance, PEG-40 Hydrogenated Castor Oil, Citric Acid, Methylisothiazolinone, Iodopropynyl Butylcarbamate, FD&C Yellow No.5, FD&C Blue No.1



GLOBAL CARE ANTIBACTERIAL HAND ALOE VERA

benzalkonium chloride liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:76 176-155
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
WATER (UNII: 059QF0K00R)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
SODIUM COCOAMPHOACETATE (UNII: W7Q5E87674)	
IODOPROPYNYL BUTYLCARBAMATE (UNII: 603P14DHEB)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
PEG-150 DISTEARATE (UNII: 6F36Q0I0AC)	
HYDROGENATED CASTOR OIL (UNII: ZF94AP8MEY)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:76176-155-01	400 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	08/01/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	08/01/2020	

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(76176-155)

Revised: 8/2020

NINGBO LIYUAN DAILY CHEMICAL PRODUCTS CO., LTD.