

**SKIN AID MEDICS ANTIBACTERIAL ALOE VERA- benzalkonium chloride 0.13% gel  
Larry (Xiamen) Hi Tech 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.*

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**Skin-aid-medics Antibacterial Hand Soap Aloe Vera**

**Active Ingredient**

Benzalkonium Chloride 0.13%

**Purpose**

Antibacterial

**Uses**

For handwashing to decrease bacteria on the skin

**Warnings**

For external use only.

**When using this product**

When using this product. Avoid contact with eyes. In case of eye contact, rinse eyes thoroughly with water.

**Stop use and ask a doctor**

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

**Keep out of reach of children.**

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

**Directions**

Pump into hands, wet as needed. Lather vigorously for at least 20 seconds. Wash skin, rinse thoroughly and dry.

**Other Information**

-Store between 15-30C (59-86F).

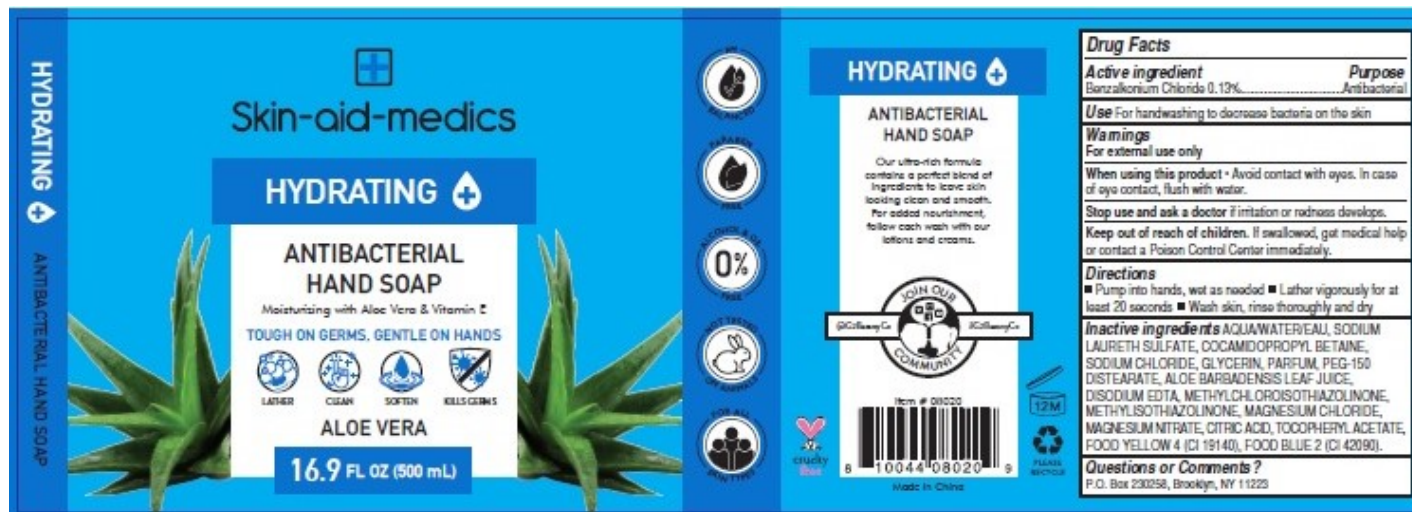
-Avoid freezing and excessive heat above 40C (104F)

**Inactive Ingredients**

Aqua Water, sodium laureth sulfate, cocoamidopropyl betaine, sodium chloride, glycerin, parfum, PEG-150 Distearate, Aloe Barbadensis Leaf Juice, Disodium EDTA, Methylchloroisothiazolinone,

Methylisothiazolinone, magnesium chloride, Magnesium Nitrate, Citric Acid, Tocopheryl Acetate, Food Yellow 4 (CI 19140) Food Blue 2 (CI 42090)

## Skin-aid-medics Antibacterial Hand Soap Aloe Vera



## SKIN AID MEDICS ANTIBACTERIAL ALOE VERA

benzalkonium chloride 0.13% gel

### Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:73166-103
Route of Administration	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.65 g in 500 mL

### Inactive Ingredients

Ingredient Name	Strength
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
MAGNESIUM NITRATE (UNII: 77CBG3UN78)	
MAGNESIUM CHLORIDE ANHYDROUS (UNII: 59XN63C8VM)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
GLYCERIN (UNII: PDC6A3C0OX)	
POLYETHYLENE GLYCOL 100000 (UNII: V46Y6OJ5QB)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
WATER (UNII: 059QF0K00R)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
SODIUM LAURETH SULFATE (UNII: BPV390UAP0)	

DISODIUM EDTA-COPPER (UNII: 6V475AX06U)

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73166-103-16	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	07/01/2020	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	07/01/2020	

**Labeler** - Larry (Xiamen) Hi Tech Co., Ltd (529759328)

**Registrant** - G2 Beauty Inc (124608169)

Revised: 8/2020

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