WET WIPES- benzalkonium chloride swab
HANGZHOU BRIGHT DAILY CHEMICAL 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.

ANTIBACTERIAL WET WIPES

Active ingredient

Benzalkonium chloride 0.1%

Purpose

Antibacterial

Use

decreases bacteria on skin

Warnings

For external use only.

Do not use

over large areas of the body

if you are allergic to any of the ingredients

when using this product,avoid contact with eyes and face.

If contact occurs,flush thoroughly with water.

Stop use and ask a doctor if irritation or rash develops and continues for more than 72 hours.

Keep out of reach of children

Keep out of reach of children. If swallowed, get medical help or contact aPoisonControlCenter right away.

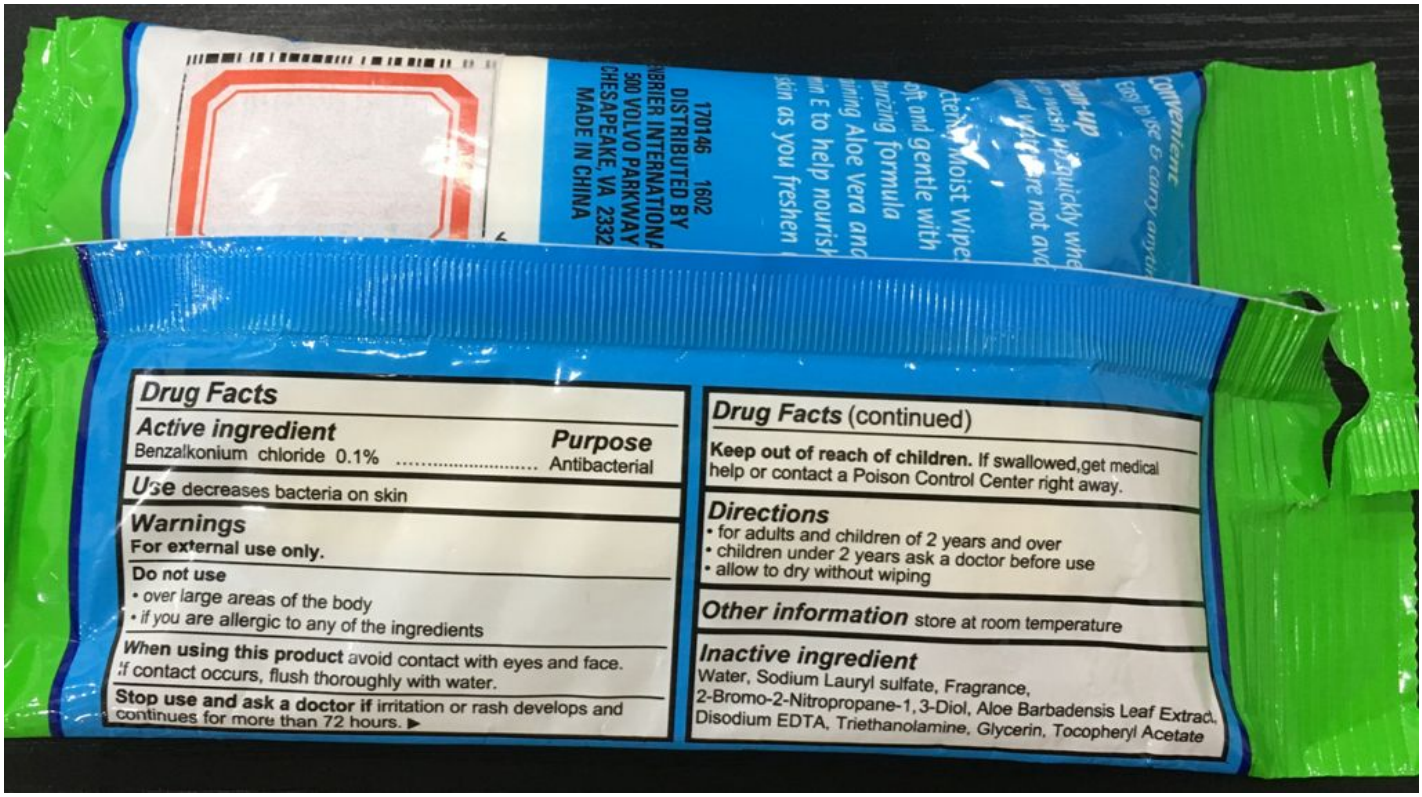
Directions

Foradults and children of 2 years and over.

children under 2 years ask a doctor before use

allow to dry without wiping

water,sodium lauryl sulfate,fragrance,2-bromo-2-nitropropane-1,3-diol,aloe barbadensis leaf extract,disodium EDTA,trithanolamine ,glycerin,tocopheryl acetate



WET WIPES

benzalkonium chloride swab

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:71198-001
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 g

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	
2-BROMO-2-NITROETHANOL (UNII: FA22WV2B2Z)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
DISODIUM EDTA-COPPER (UNII: 6V475AX06U)	
COPPER TRIETHANOLAMINE (UNII: YBM44X0B6H)	
GLYCERIN (UNII: PDC6A3C0OX)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71198-001-01	10 in 1 PACKAGE	02/13/2017	
1		0.042 g in 1 POUCH; Type 0: Not a Combination Product		
2	NDC:71198-001-02	15 in 1 PACKAGE	02/13/2017	
2		0.06 g in 1 POUCH; Type 0: Not a Combination Product		
3	NDC:71198-001-03	30 in 1 PACKAGE	02/13/2017	
3		0.126 g in 1 POUCH; Type 0: Not a Combination Product		
4	NDC:71198-001-05	80 in 1 PACKAGE	02/13/2017	
4		0.336 g in 1 POUCH; Type 0: Not a Combination Product		
5	NDC:71198-001-04	50 in 1 PACKAGE	02/13/2017	
5		0.21 g in 1 POUCH; Type 0: Not a Combination Product		
6	NDC:71198-001-06	100 in 1 PACKAGE	02/13/2017	
6		0.42 g in 1 POUCH; Type 0: Not a Combination Product		
7	NDC:71198-001-07	1200 in 1 PACKAGE	02/13/2017	
7		5.04 g in 1 POUCH; Type 0: Not a Combination Product		
8	NDC:71198-001-08	50 in 1 BOTTLE	02/13/2017	
8		0.21 g in 1 POUCH; Type 0: Not a Combination Product		
9	NDC:71198-001-09	80 in 1 BOTTLE	02/13/2017	
9		0.336 g in 1 POUCH; Type 0: Not a Combination Product		
10	NDC:71198-001-10	100 in 1 BOTTLE	02/13/2017	
10		0.42 g in 1 POUCH; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	02/13/2017	

Labeler - HANGZHOU BRIGHT DAILY CHEMICAL CO.,LTD (543255067)

Establishment

Name	Address	ID/FEI	Business Operations
HANGZHOU BRIGHT DAILY CHEMICAL CO.,LTD		543255067	manufacture(71198-001)

Revised: 1/2020 HANGZHOU BRIGHT DAILY CHEMICAL CO.,LTD