CLEANSING TOWELETTE- benzalkonium chloride swab
Tongzhou Deqi Medical Products Factory

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 Ingredients

Benzalkonium Chloride 0.13%

Purpose

Antiseptic

Use

Cleasing of face,hands and body without soap and water.Air dryies in seconds.

Non-sterile solution

Applicator is sterile if package is intact.

Warnings

For external use only
Flammable,keep away for fire or flame

Do not use:

* In large quantities
* over large areas of the body
* in eyes
* over raw or blistered areas

Stop use and ask a doctor if conditions worsen or persist for more than 7 days or clear up and occur again within a few days.

Keep out of reach of children.if swallowed get medical help or contact Poison control Center right away.

Directions

Adults and Children 2 year and older; Apply to cleaned affected area not more than 3 times daily.
Children under 2 years of age ; Consult a doctor

Inactive Ingredient

5% v/v alcohol,Purified Water

NDC: 71222-005-01

Cleansing Towelette

Drug Facts

Active Ingredients — **Purpose**
Benzalkonium chloride 0.13% . . . . . Antiseptic

Use prevent infection in minor scrapes

Non-sterile Solution
Applicator is sterile if package is intact

Drug Facts (continued)

Warnings

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Do not use: ■ in large quantities ■ over large areas of the body ■ in eyes ■ over raw or blistered areas.

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CLEANSING TOWELETTE

benzalkonium chloride swab

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:71222-005
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 mg

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
ALCOHOL (UNII: 3K9958V90M)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:71222-005-01	1 in 1 POUCH; Type 0: Not a Combination Product	05/01/2016	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	04/11/2014	

Labeler - Tongzhou Deqi Medical Products Factory (544464252)

Establishment

Name	Address	ID/FEI	Business Operations
Tongzhou Deqi Medical Products Factory		544464252	manufacture(71222-005)