BZK TOWELETTE- benzalkonium chloride swab
Henry Schein Inc.

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, 0.133% w/v

PURPOSE:

First Aid Antiseptic

Caution Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

USE: Antiseptic Cleansing of face, hands and body without soap and water. Air dries in seconds.

DO NOT USE: In the eyes or apply over large areas of the body.

STOP USE: If irritation, redness or other symptoms develop. Consult a doctor if the condition persists or gets worse.

Contains 1 Pad

DIRECTIONS: Tear open packet, unfold and use as washcloth

INACTIVE INGREDIENT: Distilled Water

PRINCIPAL DISPLAY PANEL - Pouch Label

HENRY SCHEIN®
112-5824

NDC 0404-5824-01

BZK
TOWELETTE

Contains Benzalkonium Chloride
For External Use Only
NON-STERILE

Contains
1 Pad

CE

Distributed by:

HENRY SCHEIN INC.
Melville, NY 11747 USA
EC REP

Henry Schein U.K. Holdings Ltd.
Gillingham ME8 0SB U.K.

HENRY SCHEIN®

112-5824

NDC 0404-5824-01

BZK TOWELETTE

Contains Benzalkonium Chloride
For External Use Only

NON-STERILE

TOALLITA BZK • No Estéril
Contiene cloruro de benzalconio • Destinado sólo para uso externo

SERVIETTA BZK • Non Stériles
Contient du chlorure de benzalkonium • Réservé à un usage externe

Distributed by:
Distribuido por: Distribué par:

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Melville, NY 11747 USA
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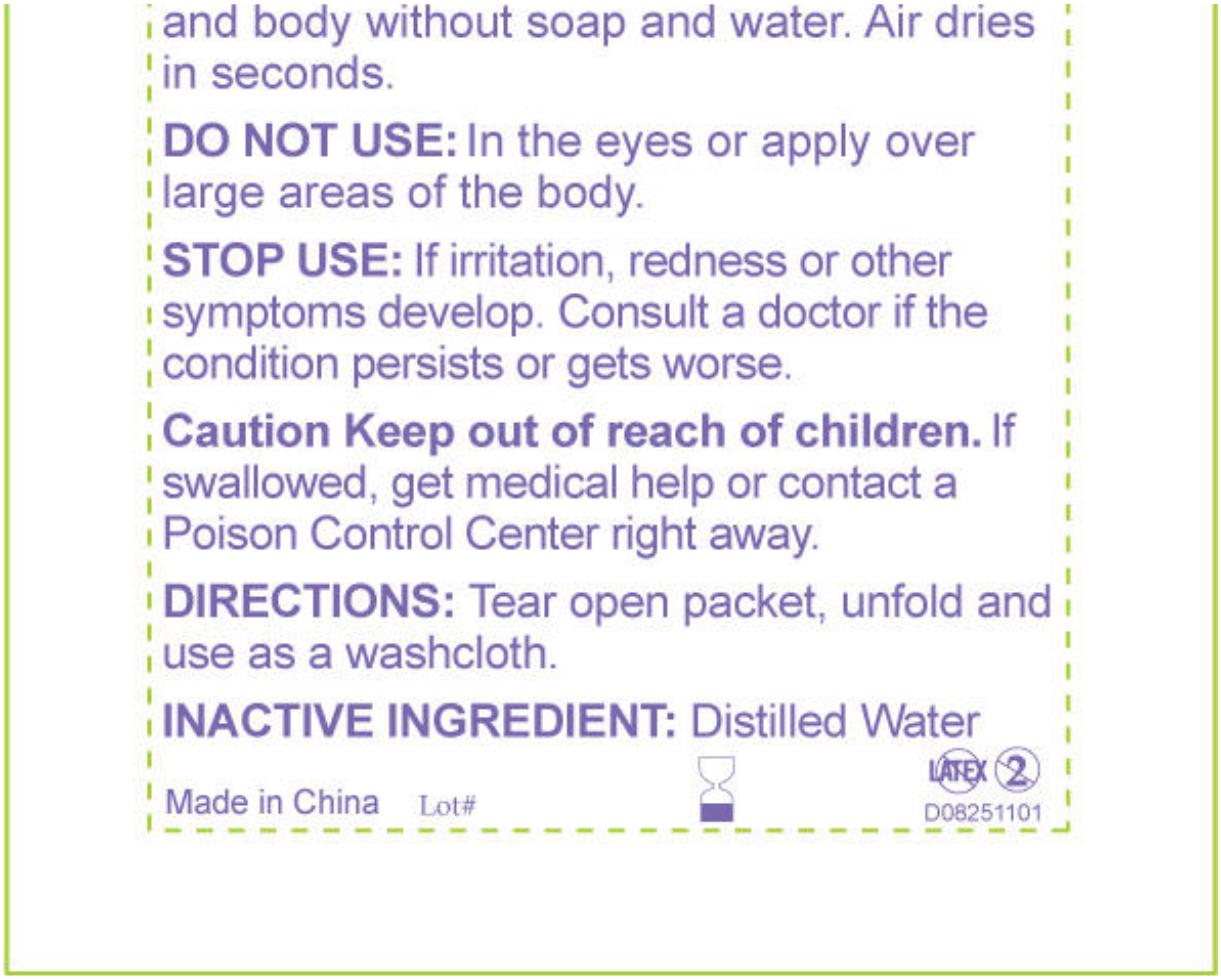

BZK TOWELETTE

benzalkonium chloride swab

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0404-5824
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM	0.001 mL in 1.4 mL

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

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0404-5824-01	1 mL in 1 POUCH		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	10/05/2011	

Labeler - Henry Schein Inc. (012430880)

Revised: 10/2011 Henry Schein Inc.