

ANTISEPTIC HAND WIPE- benzalkonium chloride patch
Taizhou Kangping Medical Science And Technology 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.

Active ingredient

Purpose - Benzalkonium Chloride 0.13% v/v.....Antiseptic

Uses

For handwashing to decrease bacteria on the skin

Warnings

For external use only.

Flammable, keep away from fire or flame.

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.

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

Directions

Tear open packet, unfold and use as a washcloth

Inactive ingredient

water

Dist. By: Total Resources International Walnut, CA 91789

www.totalresourcesintl.com



Drug Facts

Active ingredients	Purpose
Benzalkonium Chloride 0.13%.....	Antiseptic
Uses ■ Antiseptic cleansing of face, hands and body to decrease bacteria on skin without soap and water	
Warnings	
For external use only.	
Do not use in the eyes. If this happens, rinse thoroughly with water.	
Stop use and ask a doctor if irritation or redness develop and persists for more than 72 hours	
Keep out of reach of children If swallowed get medical help or contact a Poison Control Center right away	
Directions ■ tear open packet, unfold and use as a washcloth.	
Inactive ingredients	water

Dist. By: Total Resources International • Walnut, CA 91789
 ITEM #00TOW94185 Lot:20170610 Exp. 05/20

ANTISEPTIC HAND WIPE
 benzalkonium chloride patch

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:71310-556
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 g

Inactive Ingredients	
Ingredient Name	Strength
WATER (UNII: 059QF0K00R)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71310-556-01	1.6 g in 1 PACKET; Type 0: Not a Combination Product	06/01/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	06/01/2017	

Labeler - Taizhou Kangping Medical Science And Technology Co., Ltd (543429840)

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