

DIAMOND WIPES ANTIBACTERIAL HAND WIPES- benzalkonium chloride cloth
Diamond Wipes International, 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.

Diamond Wipes Antibacterial Hand Wipes

Active ingredients

Benzalkonium Chloride 0.13%

Purpose

Antibacterial

Directions

- For hand washing to decrease bacteria on skin
- recommended for repeated use

Warnings

For external use only

When using this product

avoid contact with eyes

Stop use and ask a doctor

If irritation or redness 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

- open packet
- remove and unfold wipes
- wipe hands thoroughly for approximately 15 seconds
- do not flush

Inactive ingredients

Alcohol Denat., Aloe Barbadosensis Leaf Juice, Citric Acid, Fragrance, Polysorbate 20, Sorbic Acid, Tocopheryl Acetate, Water

- Diamond Wipes
- Antibacterial Hand Wipes
- Kills 99% of Germs
- Aloe vera and Vitamin E
- Made in USA



**Kills 99%
of Germs**

1 Wipe | **Antibacterial
Hand Wipes**

Aloe Vera & Vitamin E
Cleans & Sanitizes

Made in U.S.A.

Drug Facts	
Active ingredient (in each wipe)	Purpose
Benzalkonium Chloride 0.13% w/w	Antimicrobial
Uses	
<ul style="list-style-type: none"> for hand washing to decrease bacteria on skin recommended for repeated use 	
Warnings	
<ul style="list-style-type: none"> For external use only When using this product avoid contact with eyes Stop use and contact a doctor if irritation or redness persists for more than 72 hours Keep out of reach of children. If swallowed, get medical help or contact Poison Control Center right away. 	
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Reorder Item# F1637
Distributed by
Diamond Wipes Int'l Inc.
(800) 454-1077
diamondwipes.com

DIAMOND WIPES ANTIBACTERIAL HAND WIPES
benzalkonium chloride cloth

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:64709-101
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 in 1 mL

Inactive Ingredients	
Ingredient Name	Strength
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
Sorbic acid (UNII: X045WJ989B)	
GLYCERIN (UNII: PDC6A3C0OX)	
ALCOHOL (UNII: 3K9958V90M)	
water (UNII: 059QF0KO0R)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
.ALPHA.-TOCOPHEROL ACETATE, DL- (UNII: WR1WPI7EW8)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	

Product Characteristics			
Color	white	Score	
Shape		Size	

Flavor		Imprint Code		
Contains				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:64709-101-10	1.8 mL in 1 PACKET; Type 0: Not a Combination Product	01/01/2010	
Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part333E	01/01/2010		

Labeler - Diamond Wipes International, Inc. (161104729)

Establishment			
Name	Address	ID/FEI	Business Operations
Diamond Wipes International, Inc.		161104729	manufacture(64709-101)

Revised: 2/2020

Diamond Wipes International, Inc.