

ANTIBACTERIAL WET WIPES- benzalkonium chloride swab
Zhejiang Qimei Commodity 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 0.13%

Purpose

Antibacterial

Uses

for hand washing to decrease bacteria on skin

Warnings

For External use only.

Do not use if you are allergic to any of the ingredients.

When using this product Do not get into eyes,if contact occurs ,rinse thoroughly with water.

Discontinue use if irritation or redness develops if condition persists for more than 72 hours consults a physician.

Keep out of reach of children

keep out of reach of children unless under adult supervision .if seallowed ,get medical help or contact a Poison Control Center immediately.

Directions

Thoroughly wipe hands or face with wipe,Discard in trash receptacle;do not flush.Be sure to reseal label complete to retain moisture

WATER PROPYLENE GLYCOL OCTHILINONE .ALPHA.-TOCOPHEROL

Drug Facts		Antibacterial Wet Wipes		Drug Fact (Continued)	
Active Ingredient Benzalkonium chloride 0.13%.....Antiseptic		Purpose		Directions For Use Thoroughly wipe hands or face with wipe. Discard in trash receptacle; do not flush. Be sure to reseal label complete to retain moisture.	
Uses For hand washing to decrease bacteria on skin					
Warnings <ul style="list-style-type: none"> ■ For external use only ■ Do not use if you are allergic to any of the ingredients ■ When using this products do not get into eyes, if contact occurs, rinse thoroughly with water. ■ Discontinus use if irritation or redness develops if condition persists for more than 72 hours consult a physician. ■ Keep out of reach of children unless under adult supervision. If swallowed, get medical help or contact a Poison Control center immediately. 				Inactive Ingredients Water, Propylene Glycol, Tocopheryl acetate, Polysorbate 20, Methylisothiazolinone, Methylchlorisothiazolinone	
				Made In China	

ANTIBACTERIAL WET WIPES

benzalkonium chloride swab

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69821-001
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.13 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
TOCOPHERYL RETINOATE (UNII: 0WN694NBMM)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69821-001-01	10 in 1 PACKAGE	06/01/2017	
1		0.0042 g in 1 POUCH; Type 0: Not a Combination Product		
2	NDC:69821-001-02	15 in 1 PACKAGE	06/01/2017	
2		0.0042 g in 1 POUCH; Type 0: Not a Combination Product		
3	NDC:69821-001-03	30 in 1 PACKAGE	06/01/2017	
3		0.0042 g in 1 POUCH; Type 0: Not a Combination Product		
4	NDC:69821-001-04	50 in 1 PACKAGE	06/01/2017	

4		0.0042 g in 1 POUCH; Type 0: Not a Combination Product		
5	NDC:69821-001-05	80 in 1 PACKAGE	06/01/2017	
5		0.0042 g in 1 POUCH; Type 0: Not a Combination Product		
6	NDC:69821-001-06	100 in 1 PACKAGE	06/01/2017	
6		0.0042 g in 1 POUCH; Type 0: Not a Combination Product		
7	NDC:69821-001-07	30 in 1 BOTTLE	06/01/2017	
7		0.0042 g in 1 POUCH; Type 0: Not a Combination Product		
8	NDC:69821-001-08	50 in 1 BOTTLE	06/01/2017	
8		0.0042 g in 1 POUCH; Type 0: Not a Combination Product		
9	NDC:69821-001-09	80 in 1 BOTTLE	06/01/2017	
9		0.0042 g in 1 POUCH; Type 0: Not a Combination Product		
10	NDC:69821-001-10	100 in 1 BOTTLE	06/01/2017	
10		0.0042 g in 1 POUCH; Type 0: Not a Combination Product		
11	NDC:69821-001-11	1 in 1 BAG	06/01/2017	
11		0.0042 g in 1 POUCH; Type 0: Not a Combination Product		
12	NDC:69821-001-12	250 in 1 CANISTER	06/01/2017	
12		0.0042 g in 1 POUCH; Type 0: Not a Combination Product		
13	NDC:69821-001-13	800 in 1 BAG	06/01/2017	
13		0.0042 g in 1 POUCH; Type 0: Not a Combination Product		
14	NDC:69821-001-14	1000 in 1 BAG	06/01/2017	
14		0.0042 g in 1 POUCH; Type 0: Not a Combination Product		
15	NDC:69821-001-15	1200 in 1 BAG	06/01/2017	
15		0.0042 g in 1 POUCH; Type 0: Not a Combination Product		
16	NDC:69821-001-16	1500 in 1 BAG	06/01/2017	
16		0.0042 g in 1 POUCH; Type 0: Not a Combination Product		
17	NDC:69821-001-17	16 in 1 BAG	06/01/2017	
17		0.0042 g in 1 POUCH; Type 0: Not a Combination Product		
18	NDC:69821-001-18	8 in 1 BAG	06/01/2017	
18		0.36 g in 1 POUCH; Type 0: Not a Combination Product		
19	NDC:69821-001-19	5 in 1 BAG	06/01/2017	
19		225 g in 1 POUCH; Type 0: Not a Combination Product		
20	NDC:69821-001-20	20 in 1 BAG	06/01/2017	
20		0.9 g in 1 POUCH; Type 0: Not a Combination Product		
21	NDC:69821-001-21	2000 in 1 BAG	06/01/2017	
21		5.096 g in 1 POUCH; Type 0: Not a Combination Product		
22	NDC:69821-001-22	1600 in 1 BAG	06/01/2017	
22		4.077 g in 1 POUCH; Type 0: Not a Combination Product		
23	NDC:69821-001-23	1800 in 1 BAG	06/01/2017	
23		4.587 g in 1 POUCH; Type 0: Not a Combination Product		
24	NDC:69821-001-24	2500 in 1 BAG	06/01/2017	
24		6.37 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	05/06/2015	

Establishment

Name	Address	ID/FEI	Business Operations
Zhejiang Qimei Commodity Co.,Ltd.		544331136	manufacture(69821-001)

Revised: 8/2019

Zhejiang Qimei Commodity Co.,Ltd.