

**WIPE OUT ANTIBACTERIAL WIPES FRESH SCENT- benzalkonium chloride cloth**  
**Xinsanyang Pharmaceutical (Xiamen) Co., Ltd.**

*Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, [click here](#).*

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**WIPE OUT Antibacterial Wipes Fresh Scent**

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Benzalkonium Chloride 0.13%

Purpose: Antiseptic

Antiseptic, WIPE OUT Antibacterial Wipes Fresh Scent

WIPE OUT Antibacterial Wipes Fresh Scent to help reduce bacteria that potentially can cause disease. For use when soap and water are not available.

For external use only. Flammable. Keep away from heat or flame

in children less than 2 months of age  
on open skin wounds

When using this product keep out of eyes, ears, and mouth. In case of contact with eyes, rinse eyes thoroughly with water.

Stop use and ask a doctor if irritation or rash occurs. These may be signs of a serious condition.

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

Stop use and ask a doctor if irritation or rash occurs. These may be signs of a serious condition.

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

Wipe the surface of the skin and let it dry naturally.

Supervise children under 6 years of age when using this product to avoid swallowing.

Store between 15-30C (59-86F)

Avoid freezing and excessive heat above 40C (104F)

Aloe Barbadensis Leaf Juice

Fragrance

Phenoxyethanol

Polysorbate 20

Propylene Glycol

Tocopheryl Acetata

Water



## WIPE OUT ANTIBACTERIAL WIPES FRESH SCENT

benzalkonium chloride cloth

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:80404-301
<b>Route of Administration</b>	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>BENZALKONIUM CHLORIDE</b> (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.13 in 100

### Inactive Ingredients

Ingredient Name	Strength
<b>PHENOXYETHANOL</b> (UNII: HIE492ZZ3T)	
<b>POLYSORBATE 20</b> (UNII: 7T1F30V5YH)	
<b>PROPYLENE GLYCOL</b> (UNII: 6DC9Q167V3)	
<b>.ALPHA.-TOCOPHEROL ACETATE</b> (UNII: 9E8X80D2L0)	
<b>WATER</b> (UNII: 059QF0K00R)	
<b>FRAGRANCE LAVENDER &amp; CHIA F-153480</b> (UNII: SXS9CO2TZK)	
<b>ALOE VERA LEAF</b> (UNII: ZY81Z83H0X)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:80404-301-01	1 in 1 BAG; Type 0: Not a Combination Product	09/02/2020	
2	NDC:80404-301-02	10 in 1 BAG; Type 0: Not a Combination Product	09/02/2020	
3	NDC:80404-301-03	20 in 1 BAG; Type 0: Not a Combination Product	09/02/2020	
4	NDC:80404-301-04	40 in 1 BAG; Type 0: Not a Combination Product	09/02/2020	
5	NDC:80404-301-05	60 in 1 BAG; Type 0: Not a Combination Product	09/02/2020	
6	NDC:80404-301-06	80 in 1 BAG; Type 0: Not a Combination Product	09/02/2020	
7	NDC:80404-301-07	100 in 1 BAG; Type 0: Not a Combination Product	09/02/2020	
8	NDC:80404-301-08	120 in 1 BAG; Type 0: Not a Combination Product	09/02/2020	

9	NDC:80404-301-09	200 in 1 BAG; Type 0: Not a Combination Product	09/02/2020	
10	NDC:80404-301-10	400 in 1 BAG; Type 0: Not a Combination Product	09/02/2020	
11	NDC:80404-301-11	500 in 1 BAG; Type 0: Not a Combination Product	09/02/2020	
12	NDC:80404-301-12	600 in 1 BAG; Type 0: Not a Combination Product	09/02/2020	
13	NDC:80404-301-13	800 in 1 BAG; Type 0: Not a Combination Product	09/02/2020	
14	NDC:80404-301-14	1000 in 1 BAG; Type 0: Not a Combination Product	09/02/2020	

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		09/02/2020	

**Labeler** - Xinsanyang Pharmaceutical (Xiamen) Co., Ltd. (546457554)

## Establishment

Name	Address	ID/FEI	Business Operations
Xinsanyang Pharmaceutical (Xiamen) Co., Ltd.		546457554	manufacture(80404-301)

Revised: 11/2020

Xinsanyang Pharmaceutical (Xiamen) Co., Ltd.