

SIMPLIFY ANTIBACTERIAL HAND WIPES- benzalkonium chloride cloth
Zhejiang Qimei Cosmetics Co., Ltd.

Zhejiang - Wipe (20 count)

Drug Facts

Active Ingredient

Benzalkonium chloride 0.13%

Purpose

Antiseptic

Use

to decrease bacteria on the skin

Warnings

For external use only-hands

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

When using this product

- Keep out of eyes. In case of contact with eyes, flush thoroughly with water.
- avoid contact with broken skin

Stop use and ask a doctor if

- irritation or redness develops
- condition persists for more than 72 hours

Drug Facts (continued)

Directions

- wipe hands thoroughly with product and allow to dry
- for children under 6, use only under adult supervision
- not recommended for infants

Inactive Ingredients

water (aqua), glycerin, sodium benzoate, potassium sorbate, aloe barabadsensis leaf juice, fragrance (parfum), caprylyl/capryl glucoside, polyglyceryl-4 laurate/sebacate, polyglyceryl-6 caprylate/caprinate, citric acid, polysorbate 20

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****Testing based on a subset of FDA proposal published June 17, 1994.***

Simplify ®

Antibacterial

Hand Wipes

Gentle Cleansing

- With aloe
- Scented
- Kills 99.9% of

Most Common Germs*

- Dermatologist Tested

20 Hand Wipes

OPEN

Do Not Flush

5.5 x 7 in (14 x 17.8 cm)

Simplify.

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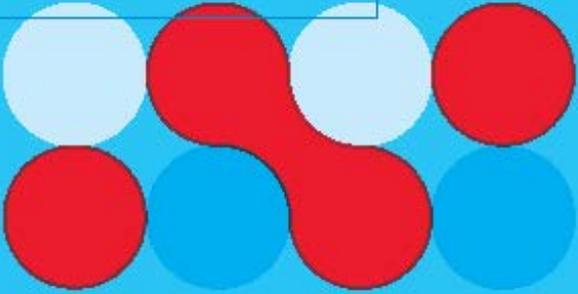
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Do Not Flush

5.5 x 7 IN (14 x 17.8 cm)

SIMPLIFY ANTIBACTERIAL HAND WIPES
benzalkonium chloride cloth

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:81773-015
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

Inactive Ingredients	
Ingredient Name	Strength
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
CAPRYLYL/CAPRYL OLIGOGLUCOSIDE (UNII: E00JL9G9K0)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
POLYGLYCERYL-4 LAURATE (UNII: 82V7NG3DYT)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
POLYGLYCERYL-6 CAPRYLATE (UNII: DGV8R54VG7)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:81773-015-01	20 in 1 POUCH; Type 0: Not a Combination Product	09/11/2023	07/01/2026

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	09/11/2023	07/01/2026

Labeler - Zhejiang Qimei Cosmetics Co., Ltd. (709887693)

Revised: 12/2025

Zhejiang Qimei Cosmetics Co., Ltd.