ANTIBACTERIAL WIPES- benzalkonium chloride swab ERC Acquisition, Inc

Drug Facts

Active ingredient

Benzalkonium chloride 0.13%

Purpose

Antimicrobial

Uses

- For hand sanitizing to decrease bacteria on the skin
- Recommended for repeated use

Warnings

For external use only

Do not use if

you are allergic to any of this product

When using this product

avoid contact with eyes. In case of eye contact, flush eyes with water.

Stop use and ask a doctor

If irritation or redness develops, or if condition persists for more than 72 hours.

Keep out of reach of children

If swallowed, get medical help or contact a Poison Control Center immediately.

Directions

Adults and Children 2 years and over: apply to hands and face. Allow to dry without wiping.

Children under 2 years: ask a doctor before use

To dispense: pull tab and remove large cover. Insert triangular threading loop through slits in cover. Do not push finger through the opening. Thread first sheet in center or roll through loop. Replace cover. Pull loop back through opening. Pull each sheet up and slightly to the side. Dispose of wipe in trash. **Do not flush**.

Other information

Store below 95 °F (35 °C). keep closed tightly. May discolor certain fabric or surfaces

Inactive ingredients

Water, SD Alcohol 40, Phenoxyethanol, Decyl Glucoside, Potassium Sorbate, Sodium Benzoate, Disodium EDTA, Citric Acid, Aloe Barbadensis (Aloe) Leaf Extract, Fragrance.

Principal Display Panel

NDC 71293-020-01

2000 count

ERC

ERC WIPING PRODUCTS

KILLS 99.9% OF GERMS

ANTIBACTERIAL WIPES

Cleans, Disinfects and Deodorizes

2000 Wipes per roll

- safe on equipment
- safe on hands
- presaturated
- made in USA

ercwipe.com 800-225-9473



ANTIBACTERIAL **WIPES**

Cleans, Disinfects and Deodorizes 2000 Wipes per Roll









ercwipe.com • 800-225-9473



Drug Facts

Active ingredient

Purpose

in 100 g

Benzalkonium chloride 0.13% For hand sanitizing to decrease bacteria on the skin Recommended for repeated use

Warnings
For external use only.

Do not use if you are allergic to any of these ingredients, When using this product, avoid contact with eyes. In case of

eve contact, flush eves with water. Stop use and ask a doctor if irritation or redness develops, or it

condition persists for more than 72 hours.

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

Directions

Adults and children 2 years and over: apply to hands and face.

Allow to dry without wiping.
Children under 2 years: ask a doctor before use.

To dispense: pull tab and remove large cover, Insert triangular threading loop through slits in cover. Do not push finger through opening. Thread first sheet in center of roll through loop. Replace cover Pull loop back through opening. Pull each sheet up and slightly to the side. Dispose of wipe in trash. Do not flush.

Other Information

Store below 95°F (35°C). Keep closed tightly. May discolor certain fabrics or surfaces.

Inactive ingredients

CHLORIDE

Water, SD Alcohol 40, Phenoxyethanol, Decyl Glucoside stassium Sorbate, Sodium Benzoate, Disodium EDTA, Citric cid, Aloe Barbadensis (Aloe) Leaf Extract, Fragrance,

Manufactured for:

ERC Wiping Products 19 Bennett Street Lynn, MA 01905

ANTIBACTERIAL WIPES

benzalkonium chloride swab

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:71293-020

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name Basis of Strength Strength BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM -**BENZ ALKONIUM** 0.13 g

UNII:7N6JUD5X6Y)

Inactive Ingredients Ingredient Name Strength WATER (UNII: 059QF0KO0R) PHENOXYETHANOL (UNII: HIE492ZZ3T) **DECYL GLUCOSIDE (UNII: Z17H97EA6Y)** POTASSIUM SORBATE (UNII: 1VPU26JZZ4)

SODIUM BENZOATE (UNII: OJ245FE5EU)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	

P	Packaging						
#	tem Code	Package Description	Marketing Start Date	Marketing End Date			
1	NDC:71293-020- 01	2000 in 1 BAG	03/16/2017				
1		1 g in 1 PACKET; Type 0: Not a Combination Product					

Marketing Information						
Marketing Application Number or Monograph Category Citation		Marketing Start Date	Marketing End Date			
OTC Monograph Drug	M003	03/16/2017				

Labeler - ERC Acquisition, Inc (019312339)

Registrant - ERC Acquisition, Inc (019312339)

Establishment					
Name	Address	ID/FEI	Business Operations		
Precare Corp.		117111327	manufacture(71293-020)		

Revised: 12/2024 ERC Acquisition, Inc