ANTIBACTERIAL WIPES- benzalkonium chloride swab
ERC Acquisition, 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.

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 ºF (35 ºC). 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

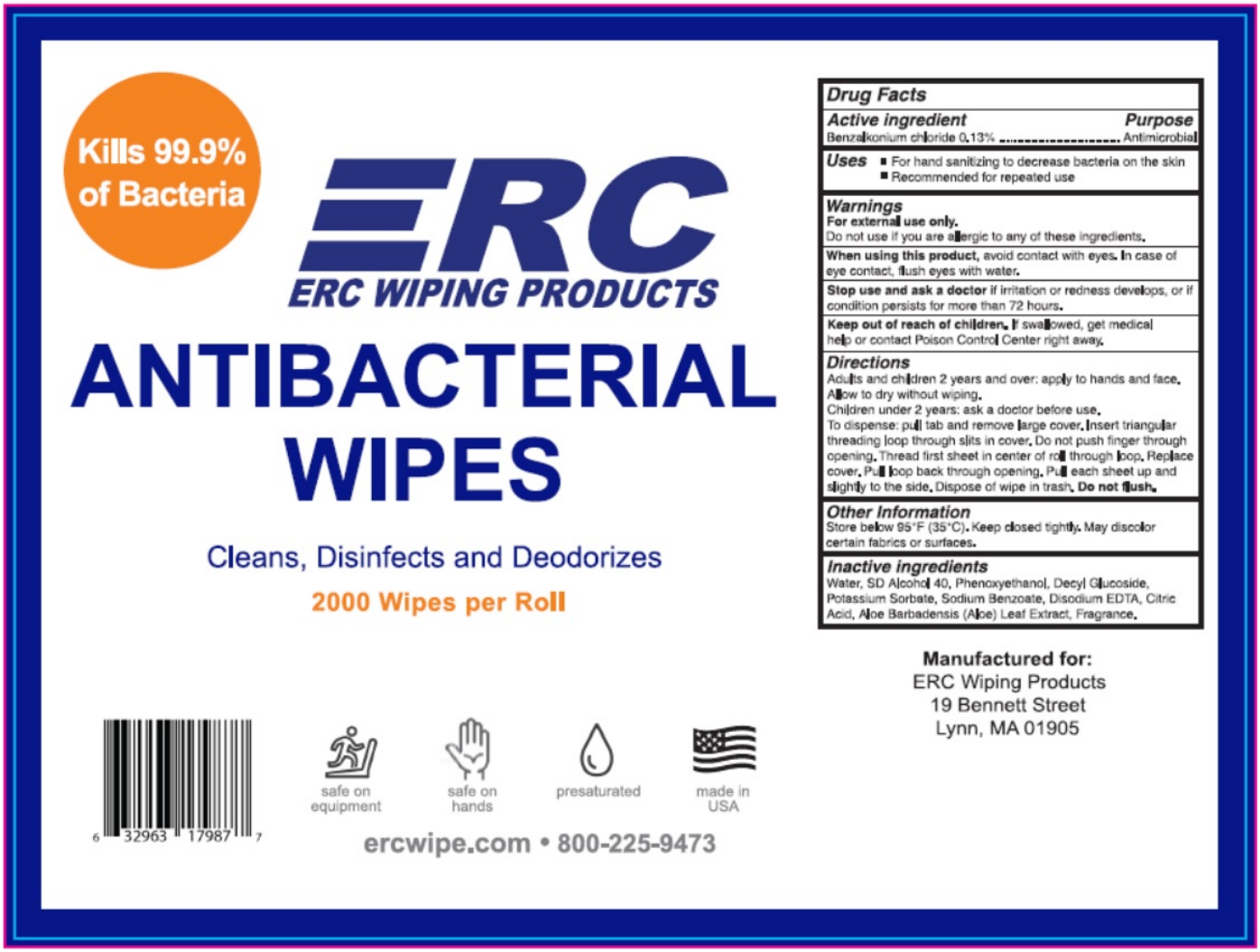
ANTIBACTERIAL WIPES

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

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 - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.13 g in 100 g

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)	

Packaging

#	Item 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 Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	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: 11/2022 ERC Acquisition, Inc