TRIP WIPES- benzalkonium chloride cloth
DETROIT WICK

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

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 right away.

Directions

- take wipe and rub thoroughly over all surfaces of both hands. Wet hands thoroughly with wipe
- rub hands together briskly to dry without wiping
- dispose of wipe
- do not flush

Other information

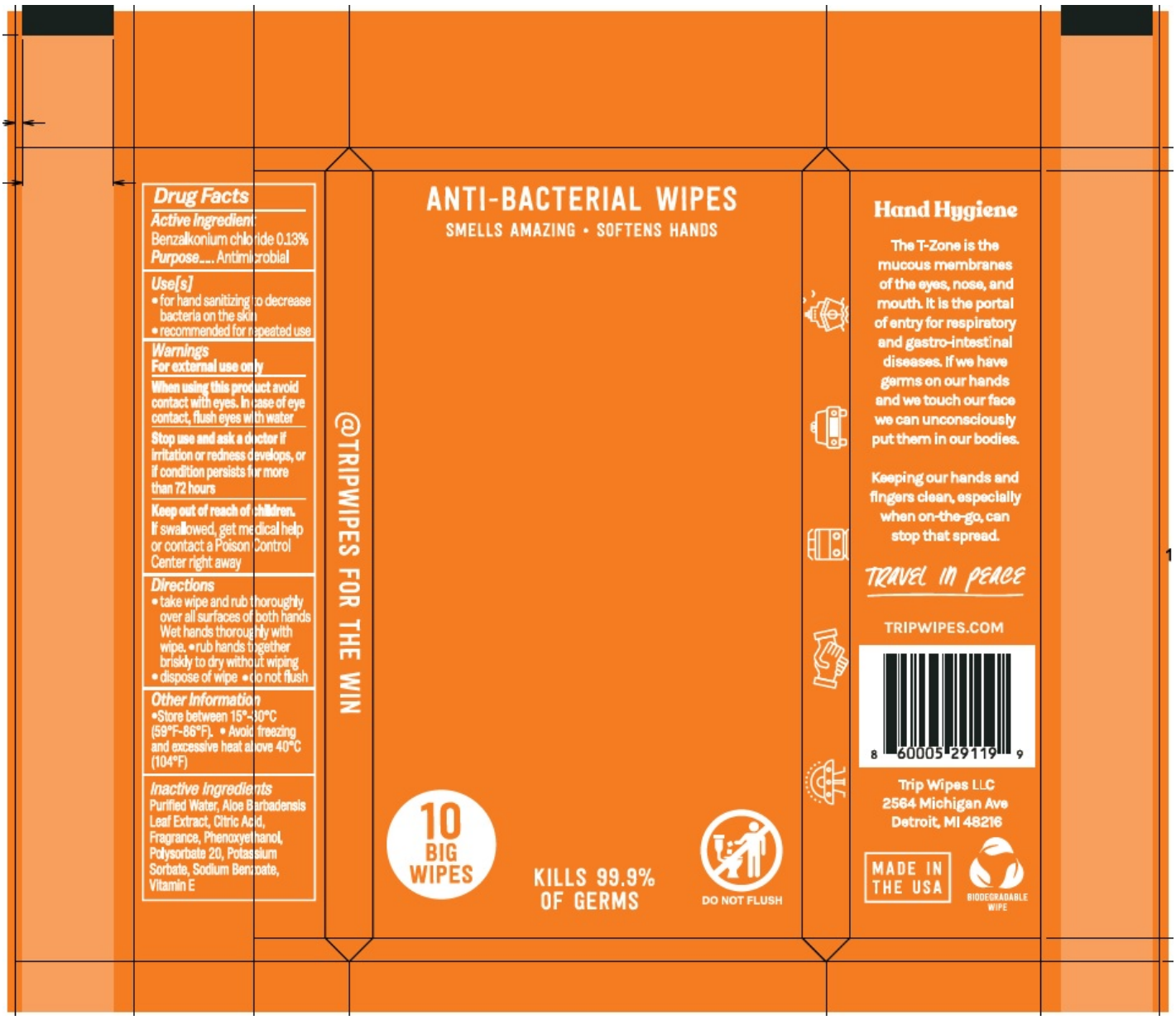
Store between 15°-30°C (59°F - 86°F).

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

Inactive ingredients

Purified Water, Aloe Barbadensis Leaf Extract, Citric Acid, Fragrance, Phenoxyethanol, Polysorbate 20, Potassium Sorbate, Sodium Benzoate, Vitamin E

Principal Display Panel



73030-025-01

ANTI-BACTERIAL WIPES

SMELLS AMAZING • SOFTENS HANDS

10 BIG WIPES

KILLS 99.9% OF GERMS

DO NOT FLUSH

TRIP WIPES

benzalkonium chloride cloth

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:73030-025
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
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
.ALPHA.-TOCOPHEROL (UNII: H4N855PNZ1)	
WATER (UNII: 059QF0KO0R)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73030-025-01	10 in 1 BOX	12/13/2022	
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	12/13/2022	

Labeler - DETROIT WICK (061117661)

Registrant - Precare Corp (858442403)

Establishment

Name	Address	ID/FEI	Business Operations
Precare Corp		858442403	manufacture(73030-025)

Revised: 12/2022

DETROIT WICK