

**TRIP WIPES- benzalkonium chloride swab
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 in a cool dry place

Inactive ingredients

Purified Water, Phenoxyethanol, Potassium Sorbate, Sodium Benzoate, Citric Acid, Polysorbate 20, Citrus Aurantium Dulcis Oil, Citrus Grandis Peel Oil, Citrus Aurantium Bergamia Fruit Oil, Elettaria Cardamomum Miniscula Seed, Linolool, Ehylene Brassylate, Citral, Kumquat Base, Galaxolide

Principal Display Panel



NDC 73030-020-02

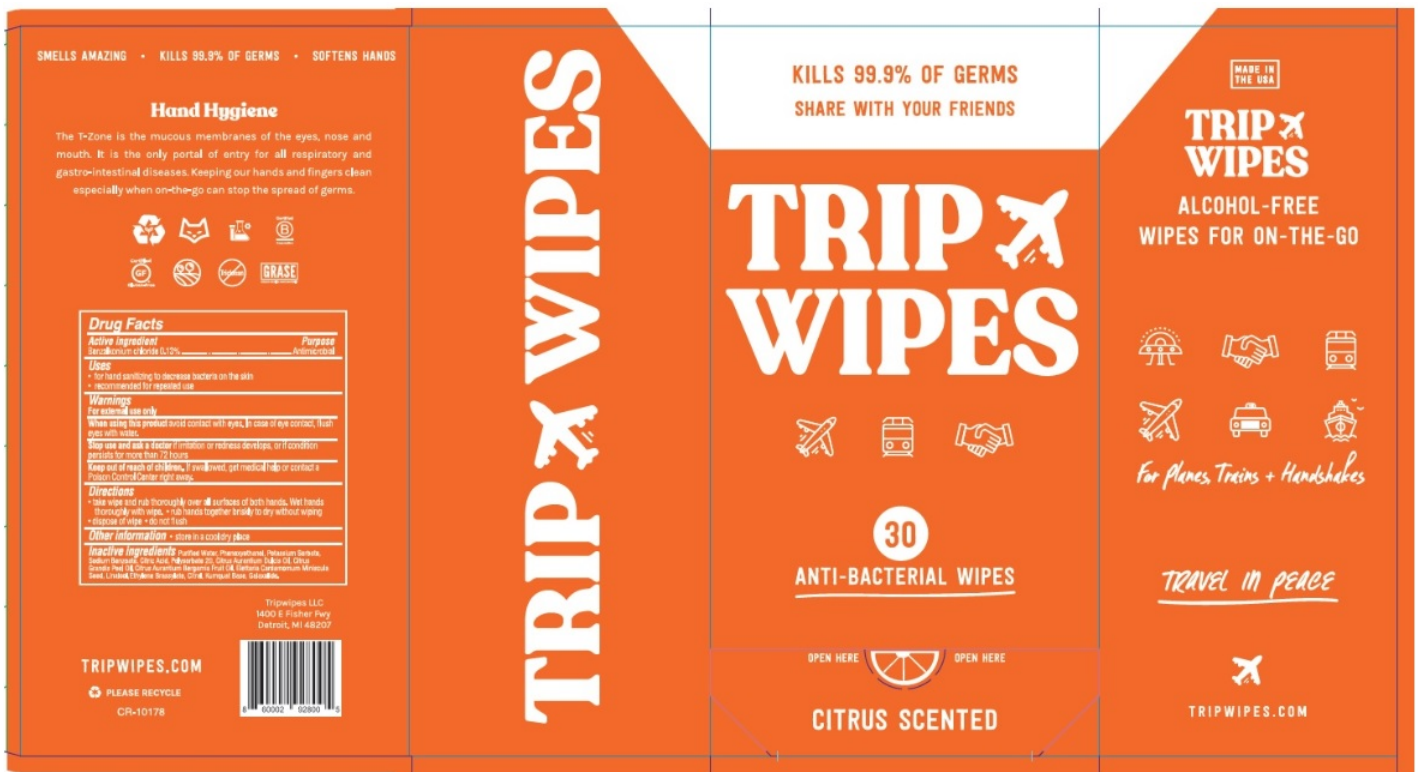
DO NOT FLUSH

CLEAN IS A WIPE AWAY

TRIP WIPES

CITRUS ANTI-BACTERIAL WIPE

1 INDIVIDUAL WIPE



NDC 73030-020-01

MADE IN THE USA

ALCOHOL-FREE WIPES FOR ON-THE-GO

For Planes, Trains +
Handshakes

TRAVEL IN PEACE

KILLS 99.9% OF GERMS

SMELLS AMAZING

SOFTENS HANDS

SHARE WITH YOU FRIENDS

30 ANTIBACTERIAL WIPES

CITRUS SCENTED

TRIPWIPES.COM

TRIP WIPES

benzalkonium chloride swab

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:73030-020
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)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
ORANGE OIL (UNII: AKN3KSD11B)	
CITRUS MAXIMA FRUIT RIND OIL (UNII: 8U3877WD44)	
BERGAMOT OIL (UNII: 39W1PKE3JI)	
LINALOOL, (+/-)- (UNII: D81QY6I88E)	
ETHYLENE BRASSYLATE (UNII: 9A87HC7ROD)	
CITRAL (UNII: T7EU009VPP)	
CITRUS JAPONICA FRUIT (UNII: 8PS197OFKT)	
HEXAMETHYLINDANOPYRAN (UNII: 14170060AT)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73030-020-01	30 in 1 BOX	04/15/2019	
1		1 g in 1 PACKET; Type 0: Not a Combination Product		
2	NDC:73030-020-02	1 in 1 PACKET	04/15/2019	
2		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	04/15/2019	

Labeler - DETROIT WICK (061117661)**Registrant** - Precare Corp (858442403)**Establishment**

Name	Address	ID/FEI	Business Operations
Precare Corp		858442403	manufacture(73030-020)

Revised: 11/2022

DETROIT WCK