HAND SANITIZING WIPES- benzalkonium chloride (0.13%) cloth
Bar-D Company

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.

Directions

- Peel back front sticker and remove wipe(s). Reseal pouch by pressing sticker back in place.
- Rub all surfaces of hands thoroughly with wipe. Allow hands to dry without additional wiping.
- Dispose of wipe in trash or compost. Do not flush.

Warnings

- For external use only.
- Avoid contact with eyes. If contact occurs, flush eyes with water.
- Stop use and ask a doctor if irritation or redness develops and persists for more than 72 hours.
- Keep out of reach of children. If swallowed, get medical help or contact a poison control center immediately

Uses

For hand sanitizing to decrase germs on skin.

Keep out of reach of children. If swallowed, get medical help or contact a poison control center immediately.

Keep out of reach of children. If swallowed, get medical help or contact a poison control center immediately.

Purpose

Antimicrobial

Active Ingredients

Benzalkonium Chloride 0.13%

Purpose: Antimicrobial

Directions

Stop use and ask doctor if irritation or redness develops and persists for more than 72

hours.

Other Information

- Store in a cool dry place.
- Keep closed tightly to retain moisture.

Inactive Ingredients

Water, Propylene Glycol, Glycerin, Phenoxyethanol, Potassium Sorbate, Disodium Cocoamphodiacetate, PEG-40 Hydrogenated Castor Oil, Aloe Vera Extract (Aloe Barbadensis), Tea Tree Oil (Melaleuca Alternifolia Leaf), Chamomile Extract (Chamomilla Recutita), Tocopheryl Acetate (Vitamin E), Peppermint Oil (Mentha Piperita), Citric Acid, Trisodium EDTA

Hand Sanitizing Wipes

HAND SANITIZING WIPES

benzalkonium chloride (0.13%) cloth

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:82303-001
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 mL

Inactive Ingredients	
Ingredient Name	**Strength**
ALPHA-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	

CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	
POLYOXYL 40 HYDROGENATED CASTOR OIL (UNII: 7YC686GQ8F)	
GLYCERIN (UNII: PDC6A3C0OX)	
TEA TREE OIL (UNII: VIF565UC2G)	
WATER (UNII: 059QF0KO0R)	
DISODIUM COCOAMPHODIACETATE (UNII: 18L9G3U51M)	
PEPPERMINT OIL (UNII: AV092KU4JH)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
CHAMOMILE (UNII: FGL3685T2X)	
EDETATE TRISODIUM (UNII: 420IP921MB)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:82303-001-02	6 in 1 BOX	12/01/2021	12/14/2023
1	NDC:82303-001-01	10 mL in 1 BAG; 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	10/01/2021	

Labeler - Bar-D Company (074734472)

Registrant - Zhejiang Qimei Commodity Co.,Ltd. (544331136)

Establishment

Name	Address	ID/FEI	Business Operations
Zhejiang Qimei Commodity Co.,Ltd.		544331136	manufacture(82303-001)

Revised: 11/2022 Bar-D Company