JANITEX ANTIBACTERIAL WET WIPES- benzalkonium chloride cloth
Pacific Health Systems 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.

***JANITEX* Antibacterial Wet Wipes**

Drug Facts

Active Ingredients

Benzalkonium Chloride (0.1%)

Purpose

Antiseptic

Uses

- To decrease bacteria on the skin that could cause disease

Warnings

- **For external use only,** harmful if swallowed.
- **Do not use** as baby wipe.
- **Do not flush.**
- **Do not use** on open skin wounds.
- **Do not use** in or near the eyes. In case of contact, rinse eyes thoroughly with water.
- **Stop use and ask a doctor** if irritation or allergic reaction occurs.
- **Flammable - Keep away from fire or flame**
- **Keep out of reach of children.** If swallowed, get medical help or contact a Poison Control Center right away.

Directions

• Open lid, gently peel back resealable label, remove and use wipes as required. Keep lid closed to prevent moisture loss.

• No need to rinse after usage. Allow skin to dry without wiping. • For adults and children 2 years and over. Discard properly after use. • Supervise children under 6 years of age when using this product to avoid swallowing.

Other Information

- Store in dry and cool place and away from oxidizers.

Inactive ingredients

Aqua, Phenoxyethanol, Propylene Glycol, Parfum, Cocamidopropyl Betaine, PEG-40 Hydrogenated Castor Oil, Benzoic Acid, Tetrasodium EDTA, Dehydroacetic Acid, Sodium Hydroxide

pH 5.5

E Vitamin

DERMATOLOGICALLY TESTED

+ moisturizer

non-woven fabric

Bleach Free

Ultra Protection

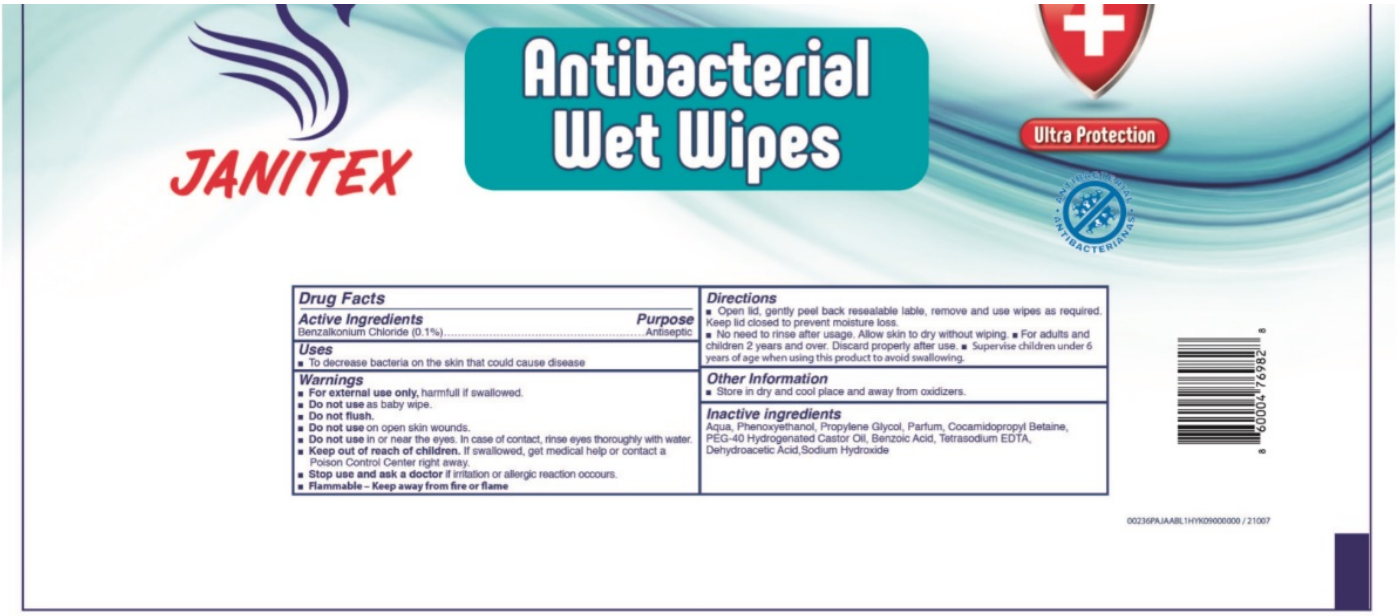
ANTIBACTERIAL

Distributed by Pacific Health Systems Inc.

19 Commerce Road Unit E Fairfield, NJ 07004

info@janitexus.com **www.janitexus.com**

Packaging



DRUG FACTS LABEL

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JANITEX ANTIBACTERIAL WET WIPES

benzalkonium chloride cloth

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:79519-050
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.1 g in 100 g

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
POLYOXYL 40 HYDROGENATED CASTOR OIL (UNII: 7YC686GQ8F)	

BENZOIC ACID (UNII: 8SKN0B0MIM)	
EDETATE SODIUM (UNII: MP1J8420LU)	
DEHYDROACETIC ACID (UNII: 2KAG279R6R)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:79519-050-01	90 in 1 PACKET	08/11/2020	
1		3.23 g in 1 PACKAGE; 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	08/11/2020	

Labeler - Pacific Health Systems Inc. (080638960)

Revised: 8/2020 Pacific Health Systems Inc.