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 Ingredient

Benzalkonium Chloride 0.1 %

Purpose

Antiseptic Hand Cleanser

Uses

Kills harmful germs

Warnings

- For external use only. Harmful if swallowed.
- When using this product, avoid contact with the eyes. If contact occurs, rinse thoroughly with water.
- Stop use and ask a doctor if irritation develops.
- Keep out of reach of children. If swallowed, call a poison control center or get medical help right away.

Directions

- Adults and children over 2 years old
- For occasional and personal domestic use
- Supervise children when they use this product
- Rub thoroughly into hands for at least 30 seconds. Allow to dry

Inactive Ingredients

Aqua, Sodium Benzoate, Phenoxyethanol, Parfum, Benzoic Acid, Dehydroacetic Acid, Panthenol, Sodium Cocoamphoacetate, Sodium Lauriminodipropionate, Octyldodecanol

Other Informations

Store in a cool & dry place

Questions?

+1 (862) 318-4763

ANTIBACTERIAL

Ultra Protection

pH 5.5

E Vitamin

DERMATOLOGICALLY TESTED

+ Moisturizer

Non-Woven Fabric

Bleach Free

Distributed by Pacific Health Systems Inc.

19 Commerce Road Unit E Fairfield, NJ 07004

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STORAGE AND DISPOSAL

- Do not leave canisters open during storage
- Do not reuse wipe
- Dispose of used wipe in trash
- Do not reuse or refill canister. Offer for recycling if available

Made in Turkey

Packaging



DRUG FACTS LABEL



JANITEX ANTIBACTERIAL WET WIPES

benzalkonium chloride cloth

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Product	Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:79519-021

Route of Administration TOPICAL

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

Inactive Ingredients			
Ingredient Name	Strength		
WATER (UNII: 059QF0KO0R)			
SODIUM BENZOATE (UNII: OJ245FE5EU)			
PHENOXYETHANOL (UNII: HIE492ZZ3T)			
BENZOIC ACID (UNII: 8 SKN0 B0 MIM)			
DEHYDRO ACETIC ACID (UNII: 2KAG279R6R)			
PANTHENOL (UNII: WV9CM0O67Z)			
SODIUM CO CO AMPHO ACETATE (UNII: W7Q5E87674)			
SO DIUM LAURIMINO DIPRO PIO NATE (UNII: 7G447D0 DH9)			
OCTYLDO DECANO L (UNII: 46 1N1O 6 14Y)			

Packaging					
# Item Code	Package Description	Marketing Start Date	Marketing End Date		
1 NDC:79519-021-01	160 in 1 CANISTER	09/18/2020			
1	3 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	09/18/2020		

Labeler - Pacific Health Systems Inc. (080638960)

Revised: 9/2020 Pacific Health Systems Inc.