NICE AND CLEAN ANTIBACTERIAL HAND WIPES- benzalkonium chloride cloth Professional Disposables International, 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.

DRUG FACTS

Use

For hand washing to decrease bacteria on the skin

Warnings

- For external use only.
- If swallowed, get medical help or contact a Poison Control Center right away.
- Do not use in the eyes.
- Flammable, keep away from fire or flame.
- Discontinue use if irritation and redness develop. If condition persists for more than 72 hours consult a doctor.

Keep out of reach of children.

Directions

- Peel back label slowly.
- Pull out wipe and reseal label.
- Unfold and use; discard after single use.
- Wet hands thoroughly with product and allow to dry without wiping.

Other information

Lot No. and Exp. date - on side

Dosage

Wet hands thoroughly with product and allow to dry without wiping.

Inactive ingredients

Water, SD Alcohol 40, Sorbic Acid, PPG-2 Hydroxyethyl Cocamide, Disodium EDTA, Aloe Barbadensis Leaf Juice, Fragrance

Manufactured for: Nice-Pak Products, Inc.

Orangeburg, NY 10962-1376 - USA - 1-800-999-6423 Made in USA

Principal Display Panel - 20 Count

Nice and Clean Antibacterial Hand WipesKills 99.99% of Germs
Hypoallergenic with Aloe and Moisturizer

20 Wipes [5.0 in x 8.0 in (12.7 cm x 20.3 cm)] 12 Packs per Tray

Lid Label

resealable A



Hand Wipes

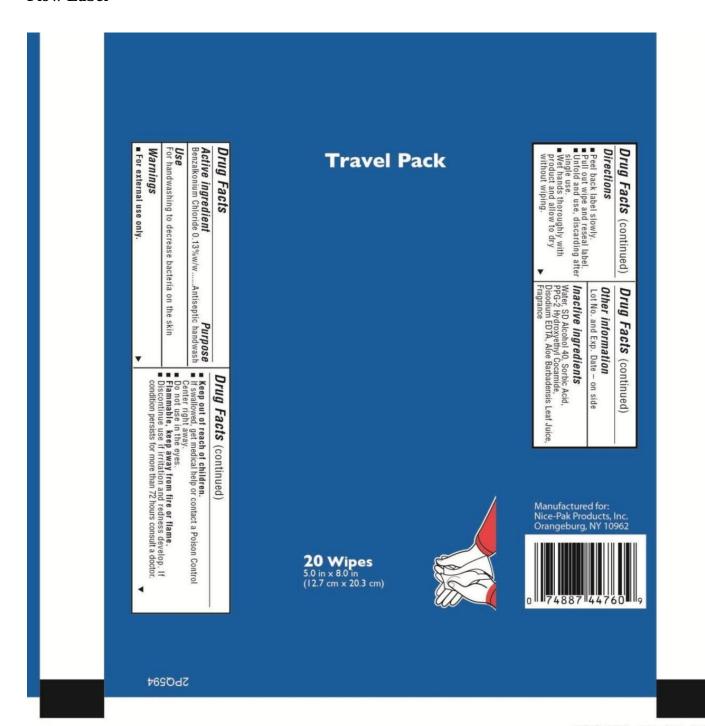


Kills 99.99% of Germs!

Hypoallergenic with Aloe & Moisturizer

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Flow Label



Tray Label



Active ingredient

Benzalkonium Chloride 0.13% w/w

Purpose

Antiseptic handwash

NICE AND CLEAN ANTIBACTERIAL HAND WIPES						
benzalkonium chloride cloth						
Product Information						
Product Type	HUMAN OTC DRUG	Item Code (Source) NDC:		NDC:1081	0819-7006	
Route of Administration	TOPICAL					
Active Ingredient/Active Moi	ety					
Ingredient Name		Basis of Strength		Strength		
Benzalkonium Chloride (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)		Benzalkonium Chloride		0.13 mg in 1 mL		

Inactive Ingredients		
Ingredient Name	Strength	
WATER (UNII: 059QF0KO0R)		
ALCOHOL (UNII: 3K9958V90M)		
SORBIC ACID (UNII: X045WJ989B)		
COCO DIETHANOLAMIDE (UNII: 92005F972D)		
EDETATE DISO DIUM (UNII: 7FLD91C86K)		
ALOE VERA LEAF (UNII: ZY8 1Z8 3H0 X)		
MICRO CITRUS AUSTRALIS FRUIT (UNII: 9 DNS80T428)		

Packaging			
# Item Code	Package Description	Marketing Start Date	Marketing End Date
1 NDC:10819-7006-1	12 in 1 TRAY		
1	38.3 mL in 1 PACKET		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	0 1/0 1/20 12	

Labeler - Professional Disposables International, Inc. (800777117)

Establishment					
Name	Address	ID/FEI	Business Operations		
Professional Disposables International, Inc.		800777117	manufacture(10819-7006)		

Revised: 3/2013 Professional Disposables International, Inc.