SANI PROFESSIONAL BRAND SANITIZING 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 immediately.
- 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
- To dispense, lift cover, remove seal, pull center sheet from roll, twist to a point, feed through dispenser hole in cover. Keep lid closed to prevent moisture loss.
- Wet hands thoroughly with product and allow to dry without wiping.
- Discard after single use.

Other information

See bag label

Dosage

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

Inactive ingredients

Water, SD Alcohol 40, Sorbic Acid, Cocamide DEA, Disodium EDTA, Aloe Barbadensis Leaf Juice, Fragrance

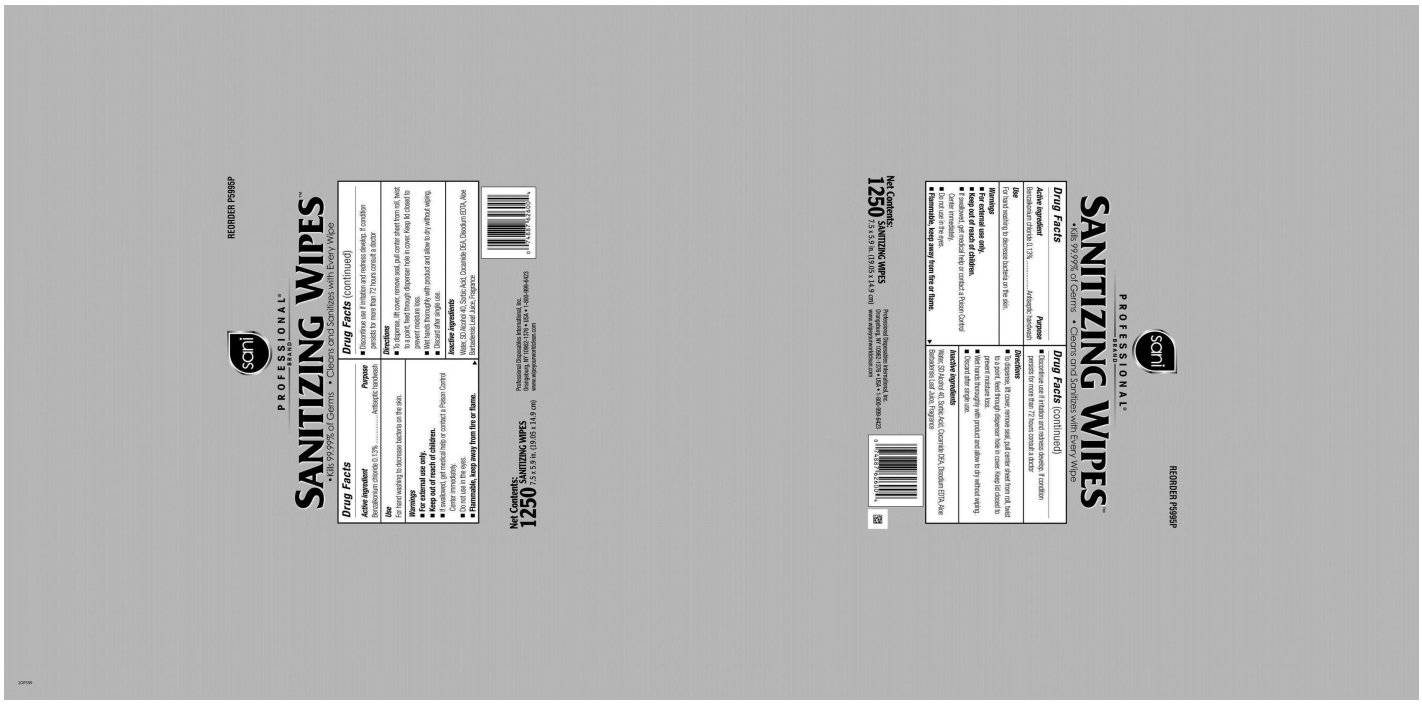
Professional Disposables International, Inc.
Orangeburg, NY 10962-1376 - USA - 1-800-999-6423
Made in USA www.wipeyourworldclean.com

Principal Display Panel - 1250 Count

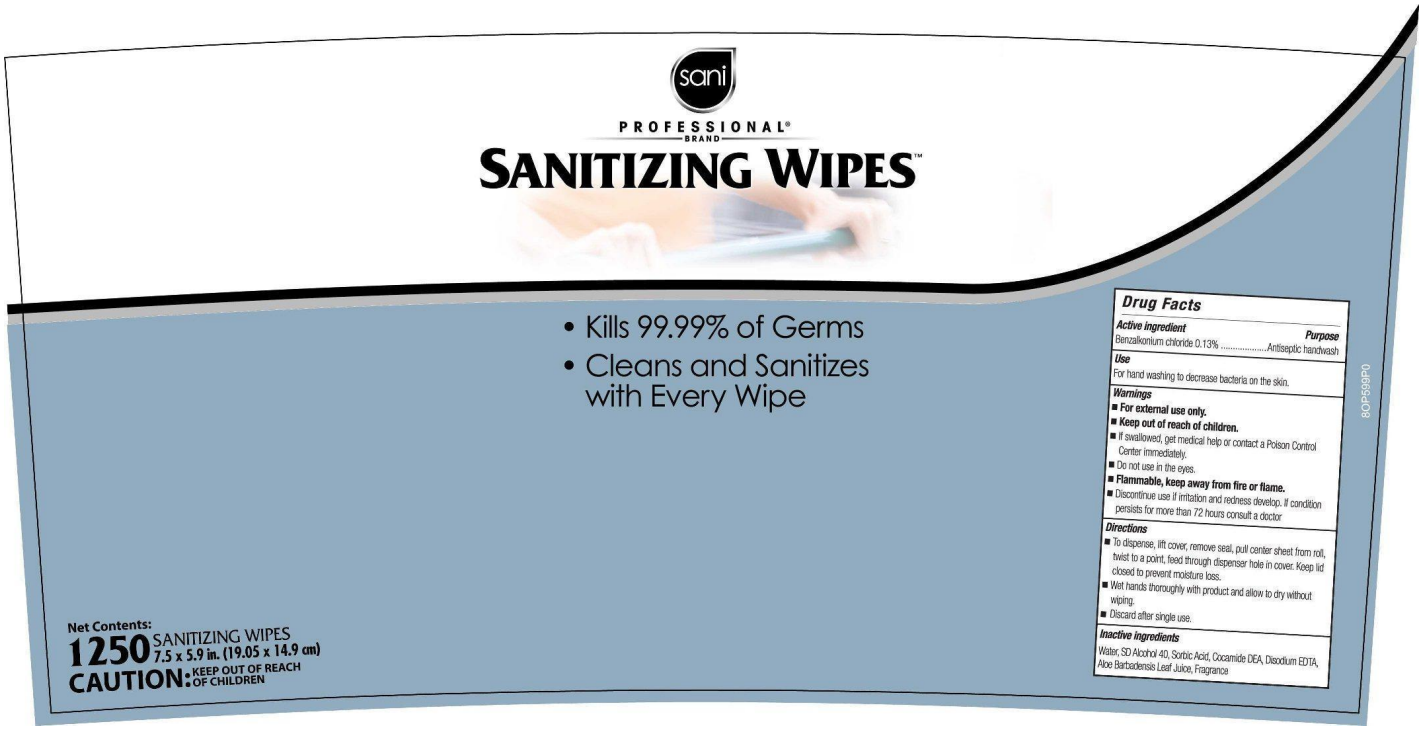
Sani Professional Brand
Sanitizing Wipes
Kills 99.99% of Germs
Cleans and Sanitizes with Every Wipe

1250 Sanitizing Wipes [7.5 x 5.9 in (19.05 x 14.9 cm)]
CAUTION: KEEP OUT OF REACH OF CHILDREN

Bag Label



Pail Label



Active ingredient

Benzalkonium Chloride 0.13% w/w

Purpose

Antiseptic handwash

SANI PROFESSIONAL BRAND SANITIZING WIPES

benzalkonium chloride cloth

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:10819-7002
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 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
ALCOHOL (UNII: 3K9958V90M)	
SORBIC ACID (UNII: X045WJ989B)	
COCO DIETHANOLAMIDE (UNII: 92005F972D)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
MICROCITRUS AUSTRALIS FRUIT (UNII: 9DNS80T428)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:10819-7002-2	1250 in 1 PAIL		
1		4674 mL in 1 BAG		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	01/01/2012	

Labeler - Professional Disposables International, Inc. (800777117)

Establishment

Name	Address	ID/FEI	Business Operations
Professional Disposables International, Inc.		800777117	manufacture(10819-7002)

Revised: 2/2013 Professional Disposables International, Inc.