PDI SANI HANDS- instant hand sanitizing wipes 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.

Sani-Hands Bedside Pack NDC 10819-3918-1

Active ingredient

Alcohol 70% by volume

Purpose

Antiseptic

Uses

- Antiseptic.
- For handwashing to decrease bacteria on the skin.
- Recommended for repeated use.
- Dries in seconds.

Warnings

- Flammable, keep away from fire or flame.
- For external use only.
- Do not use in or contact the eyes.
- Discontinue use if irritation and redness develop. If conditions persist for more than 72 hours, consult a physician.

Keep this out of reach of children

unless under adult supervision.

if swallowed, get medical help or contact a Poison Control Center immediately

Directions

- Peel back label slowly.
- Pull out wipe and reseal label.
- Unfold and use.
- Wipe hands, fingers, interdigital areas and wrists thoroughly with towelette. Be sure to utilize the entire wipe surface. Allow to dry.
- If hands are visibly soiled or contaminated, use first wipe to clean hands, then discard wipe. Sanitize with a second wipe.
- Discard after single use.

inactive ingredients

Water, Propylene Glycol, Glycerin, Aloe Barbadensis Leaf juice, Tocopheryl Acetate

Principal Display panel

Reseal lid:



Flowrap:



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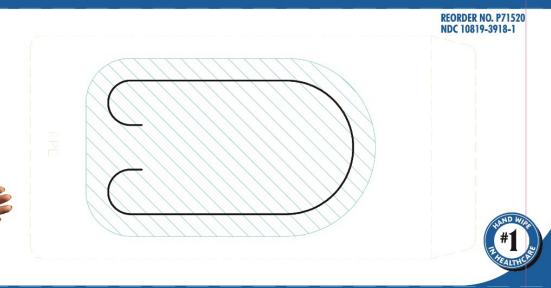
Other information

Lot number and expiration date can be found on pack.

Inactive ingredients

Water, Propylene Glycol, Glycerin, Aloe Barbadensis Leaf Juice, Tocopheryl Acetate





Kills 99.99% of Germs • Contains Moisturizing Aloe & Vitamin E

Drug Facts

Active ingredient Purpose Ethyl Alcohol 70% by volume. .. Antiseptic

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Drug Facts (continued)

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Directions

■ Peel back label slowly.
■ Pull out wipe and reseal label.
■ Unfold and use.
■ Wipe hands, fingers, interdigital areas



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instant hand sanitizing wipes cloth

Product Information

Product Type HUMAN OTC DRUG NDC:10819-3918 Item Code (Source)

TOPICAL Route of Administration

Active Ingredient/Active Moiety

	Ingredient Name	Basis of Strength	Strength
ı	ALCOHOL (UNII: 3K9958 V90M) (ALCOHOL - UNII:3K9958 V90M)	ALCOHOL	0.7 mL in 1 mL

Inactive Ingredients				
Ingredient Name	Strength			
WATER (UNII: 059QF0KO0R)				
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)				
ALOE VERA LEAF (UNII: ZY8 1Z8 3H0 X)				
ALPHA-TO COPHEROL ACETATE (UNII: 9E8X80D2L0)				
GLYCERIN (UNII: PDC6 A3C0 OX)				

Packaging	ackaging					
# Item Code	Package Description	Marketing Start Date	Marketing End Date			
1 NDC:10819-3918-	20 in 1 CELLO PACK	04/01/2016				
1	4.15 mL in 1 APPLICATOR; Type 0: Not a Combination Product					

Marketing Infor	mation		
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	04/01/2016	

Labeler - Professional Disposables International Inc (800777117)

 $\pmb{Registrant - \text{Professional Disposables International Inc (800777117)}}$

Revised: 4/2019 Professional Disposables International Inc