NURI CARE HAND SANITIZING WIPES- hand sanitizing wipes cloth SUNJU CORPORATION

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.

Active Ingredient(s)

Benzalkonium Chloride 0.1% v/v. Purpose: Antiseptic

Purpose

Antiseptic

Use(s)

Hand sanitizer to help reduce bacteria on the skin.

Warnings

Children under 2 years ask a doctor before use.

Do not use

- in children less than 2 months of age
- on open skin wounds

When using this product keep out of eyes, ears, and mouth. In case of contact with eyes, rinse eyes thoroughly with water.

Stop use and ask a doctor if irritation or rash develops and continues for more than 72 hours.

Keep out of reach of children. if swallowed, get medical help or contact a Poison Control Center right away.

Directions

- Wet hands thoroughly with product and allow to dry
- Supervise children under 6 years of age when using this product

Other information

• Store at room temperature

Inactive ingredients

Water, Sodium Hyaluronate

Package Label - Principal Display Panel



NURI CARE HAND SANITIZING WIPES									
hand sanitizing wipes cloth									
Product Informa	tion								
Product T ype	oduct Type HUMAN OTC DRUG			Item Code (Source)		NDC:75272-002			
Route of Administra	tion	TOPICAL							
Active Ingradian	t/Active Moi	0. fx7							
Active Ingredient/Active Moiety Ingredient Name Basis of						Strength			
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM -				BENZALKONIUM		0.1 g			
UNII:7N6 JUD5X6 Y)				CHLORIDE		in 100 g			
Inactive Ingredients									
Ingredient Name						Strength			
SO DIUM HYDRO XIDE (UNII: 55X04QC32I)									
WATER (UNII: 059QF0KO0R)									
Packaging									
# Item Code		Package Description Market		ing Start Date Market		ng End Date			
1 NDC:75272-002-50	002-50 470 g in 1 PACKET; Type 0: Not a Combina		Product 05/13/2020						
Marketing Information									
Marketing Catego	ry Applicat	ion Number or Monograph C	itation Mark	Marketing Start Date		Marketing End Date			
OTC monograph not fin	nal part333A	0		020					

Labeler - SUNJU CORPORATION (694816019)

Establishment							
Name	Address	ID/FEI	Business Operations				
Nuricham, Inc.		695541121	manufacture(75272-002)				

Revised: 5/2020

SUNJU CORPORATION