BATHEAL 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%

Purpose

Antibacterial

Use(s)

Hand sanitizer to help reduce bacteria on the skin.

Warnings

For External use only. For children under 3 years of age, consult with a doctor before use.

Do not use

- on large areas of the body
- if you are allergic to any of the ingredients

When using this product do not use in or near the eyes. In case of contact, 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

- For adults and children of 3 years and over, 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, Alcohol, Sodium Hydroxide, Fragnance

Package Label - Principal Display Panel













Effective at eliminating most common germs and bacteria.



smrsb to %9.9% of Germs

Made in Korea

Batheal HAND SANITIZING

WIPES

Kills 99.9% of Germs*

NDC 75272-004-10

7.9 IN X 5.9 IN (20cm X 15cm)

30 sheets

Fresh Scent

Drug Facts Active ingredient[s] Benzalkonium Chloride 0.1%

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Drug Facts (continued)

Stop use ask a doctor if irritation or rash develops and

continues for more than 72 hours.

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- For adults and children of 3 years and over, wet hands thouroughly with product and allow to dry.
 Supervise children under 6 years of age when using this product.

Inactive ingredients Water, Alcohol, Sodium Hydroxide

Manufactured by : SHINE Co., Ltd. 55-1, Meonosimi-qil, 264beon-qil, Geumwano-eu

hand sanitizing wipes cloth

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:75272-004	
Route of Administration	TOPICAL			

l	Active Ingredient/Active Moiety			
ı	Ingredient Name	Basis of Strength	Strength	
	BENZALKO NIUM CHLO RIDE (UNII: F5UM2KM3W7) (BENZALKO NIUM - UNII:7N6 JUD5X6 Y)	BENZALKONIUM CHLORIDE	0.1 g in 100 g	

Inactive Ingredients			
Ingredient Name	Strength		
ALCOHOL (UNII: 3K9958V90M)			
SO DIUM HYDRO XIDE (UNII: 55X04QC32I)			
WATER (UNII: 059QF0KO0R)			

Pac	kaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1 NI	DC:75272-004-10	108 g in 1 PACKET; Type 0: Not a Combination Product	06/08/2020	

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part333A	06/08/2020		

Labeler - SUNJU CORPORATION (694816019)

Establishment				
Name	Address	ID/FEI	Business Operations	
SHINE CO.,LTD		688437450	manufacture(75272-004)	

Revised: 6/2020 SUNJU CORPORATION