HAND BODY WIPE- hand body wipe cloth HEBEI JIATING HEALTH PRODUCT CO.,LTD

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.

80768-001 hand body wipe

Active Ingredient(s)

Benzalkonium chloride 0.02% Benzethonium chloride 0.02%

Purpose

Antiseptic

Use

Decreases bacteria on skin

Warnings

For external use only.

Do not use:

If you are allergic to any of the ingredients

When Using:

Do not get into eyes. If contact occurs. rinse thoroughly with water

O T C - Stop Use:

If irritation or rash develops and continues for more than 72 hours

OTC-KEEPOUTOFREACI-IOFCI-IILDREN

If swallowed, get medical help or contact a Poison Control Center right away

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Directions

Double layer seal.which reduces pollution and locks water. Increase the size and thickness, better for use, pearl grain design, easy to clean. The skin feels comfortable and non-sticky, giving people more refreshing and comfortable after using it. Suitable for wiping hands, mouth, face, skin and other daily necessities.

Other information

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Inactive ingredients

IPBC, Pure water, Mint essence

Package Label - Principal Display Panel

















































HAND BODY WIPE

hand body wipe cloth

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:80768-001	
Route of Administration	TOPICAL			

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
BENZETHO NIUM CHLO RIDE (UNII: PH41D05744) (BENZETHONIUM - UNII:1VU15B70BP)	BENZETHONIUM CHLORIDE	0.02 g in 100		
BENZALKO NIUM CHLO RIDE (UNII: F5UM2KM3W7) (BENZALKO NIUM - UNII:7N6 JUD5X6 Y)	BENZALKONIUM CHLORIDE	0.02 g in 100		

Inactive Ingredients			
Ingredient Name	Strength		
IODOPROPYNYL BUTYLCARBAMATE (UNII: 603P14DHEB)			
MINT (UNII: FV98Z8GITP)			
WATER (UNII: 059QF0KO0R)			

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:80768-001-01	10 in 1 BAG; Type 0: Not a Combination Product	03/30/2020	
2	NDC:80768-001-03	30 in 1 BAG; Type 0: Not a Combination Product	03/30/2020	
3	NDC:80768-001-02	20 in 1 BAG; Type 0: Not a Combination Product	03/30/2020	
4	NDC:80768-001-04	40 in 1 BAG; Type 0: Not a Combination Product	03/30/2020	
5	NDC:80768-001-05	50 in 1 BAG; Type 0: Not a Combination Product	03/30/2020	
6	NDC:80768-001-06	60 in 1 BAG; Type 0: Not a Combination Product	03/30/2020	
7	NDC:80768-001-07	70 in 1 BAG; Type 0: Not a Combination Product	03/30/2020	
8	NDC:80768-001-08	80 in 1 BAG; Type 0: Not a Combination Product	03/30/2020	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	03/30/2020	

Labeler - Hebei Jiating Health Product co.,Ltd (554530310)

Establishment				
Name	Address	ID/FEI	Business Operations	
HEBEI JIATING HEALTH PRODUCT CO.,LTD		554530310	manufacture (80768-001)	

Revised: 10/2020 HEBEI JIATING HEALTH PRODUCT CO.,LTD