FUSION ANTIBACTERIAL WIPES- antibacterial wet wipes cloth
Fusion Products 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.

Fusion Antibacterial Wipes

Active ingredient

Benzalkonium Chloride 0.13% w/w

Purpose

Antiseptic

Uses

For hand washing to decrease bacteria on the skin.

Warnings

For external use only

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 appears and lasts

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

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 appears and lasts

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

Directions

Apply to clean dry hands.

Wet hands thoroughly with the product and allow to dry

Children under 6 should be supervised when using this product

Other information

Do not store above 104°F (40°C)

Do not use if seal is missing or broken

Inactive Ingredients

aloe barbadensis leaf juice, decyl glucoside, glycerin, water.

70 WIPES, NDC: 80336-004-09

FUSION ANTIBACTERIAL WIPES

antibacterial wet wipes cloth

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:80336-004
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.0013

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
DECYL GLUCOSIDE (UNII: Z17H97EA6Y)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:80336-004-09	70 in 1 BOTTLE; Type 0: Not a Combination Product	10/23/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	10/23/2020	

Labeler - Fusion Products Ltd. (206910072)

Registrant - Hangzhou Lookon Commodity Co., Ltd. (541760483)

Establishment

Name	Address	ID/FEI	Business Operations
Hangzhou Lookon Commodity Co., Ltd.		541760483	manufacture(80336-004) , relabel(80336-004)

Revised: 6/2021 Fusion Products Ltd.