ANTIBACTERIAL HAND WASH- silver solution
Fujian Mengjiaolan Daily Chemical Co., Ltd.

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

Antibacterial Hand Wash

The Hand Wash is manufactured using only the following United States Pharmacopoeia (USP) grade ingredients in the preparation of the product (percentage in final product formulation) :

a. SILVER 0.18mg/300ml
b. SODIUM LAURETH SULFATE
c. SORBITOL
d. Sterile distilled water or boiled cold water.

The firm does not add other active or inactive ingredients. Different or additional ingredients may impact the quality and potency of the product.

Active Ingredient(s)

SILVER 0.18mg/300ml Purpose: Antiseptic

Purpose

Antiseptic, Hand Sanitizer

Use

Hand Sanitizer to help reduce bacteria that potentially can cause disease. For use when soap and water are not available.

Warnings

For external use only. Flammable. Keep away from heat or flame

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 occurs. These may be signs of a serious condition.
Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

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Directions

- Place enough product on hands to cover all surfaces. Rub hands together until dry.

- Supervise children under 6 years of age when using this product to avoid swallowing.

Other information
- Store between 15-30C (59-86F)
- Avoid freezing and excessive heat above 40C (104F)

Inactive ingredients

SODIUM LAURETH SULFATE, SORBITOL, purified water USP

Package Label - Principal Display Panel

300 mL NDC: 74531-002-01

ANTIBACTERIAL HAND WASH

silver solution

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:74531-002
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
SILVER (UNII: 3M4G523W1G) (SILVER - UNII:3M4G523W1G)	SILVER	0.18 mg in 300 mL

Inactive Ingredients	
Ingredient Name	**Strength**

SODIUM LAURETH SULFATE (UNII: BPV390UAP0)	
SORBITOL (UNII: 506T60A25R)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:74531-002-01	300 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/30/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		03/30/2020	

Labeler - Fujian Mengjiaolan Daily Chemical Co., Ltd. (546632451)

Establishment

Name	Address	ID/FEI	Business Operations
Fujian Mengjiaolan Daily Chemical Co., Ltd.		546632451	manufacture(74531-002)

Revised: 3/2020 Fujian Mengjiaolan Daily Chemical Co., Ltd.