# SUFIYA- hocl(hypochlorous acid) liquid EMSOLUTION 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.

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#### 82124-701 Sufiya

#### **Active ingredient[s]**

HOCI (Hypochlorous acid) 0.1%

#### **Purpose**

Disinfect, deodorization

#### Use[s]

Sanitizer and disinfectant 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

- for any other purpose
- in children less than 2 months of ago
- 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 any abnormality occurs on the skin.

#### Keep out of reach of children.

If swallowed, drink enough water and get medical help or contact a Poison Control Center right away.

#### **Directions**

- Spray from a distance of 4-8 inches where sterilization, disinfection, and deodorization are required
- Supervise children under 6 years of age when using this product to avoid swallowing.

#### Other information

- Store between 15-30C(58-86F)
- Avoid freezing and excessive heat above 40C(104F)
- If medical treatment is required, look at the product container or labeling information.

#### **Inactive ingredients**

Purified water

## Package Label

# Drug Facts Active ingredient[s] Purpose HOCl (Hypochlorous acid) 0.1% Disinfect, deodorization

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#### **SUFIYA**

hocl(hypochlorous acid) liquid

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:82124-701	
Route of Administration	TOPICAL			

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
HYPOCHLOROUS ACID (UNII: 712K4CDC10) (HYPOCHLOROUS ACID - UNII:712K4CDC10)	HYPOCHLOROUS ACID	0.1 g in 100 mL	

Inactive Ingredients			
Ingre	dient Name	Strength	
WATER (UNII: 059QF0KO0R)			

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:82124- 701-01	60 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	07/20/2021	
2	NDC:82124- 701-02	600 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	07/20/2021	
3	NDC:82124- 701-03	1000 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/20/2021	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		07/20/2021	
other		07/20/2021	

# Labeler - EMSOLUTION Co., Ltd (694603987)

# Registrant - EMSOLUTION Co., Ltd (694603987)

Establishment				
Name	Address	ID/FEI	Business Operations	
EMSOLUTION Co., Ltd		694603987	manufacture(82124-701)	

Revised: 12/2022 EMSOLUTION Co., Ltd