DAILY DOCTOR HYPOCHLOROUS ACID DISINFECTANT LIQUID- hypochlorous acid spray

GFC Life Science 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.

76836-002_Daily Doctor Hypochlorous acid disinfectant liquid

Active ingredient

HYPOCHLOROUS ACID 0.1%

Purpose

Antimicrobial

Use

• Prevent and treat infection of the skin

Warnings

For external use only

Do not use if sensitive of chlorine compounds.

Stop use and ask a doctor if redness, irritation or swelling or pain persists or increases.

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

Directions

• Use as liquid. Sterilize by immersing completely for about 3 to 5 minutes. For instruments that are difficult to deposit, spray and sterilize for about 3 minutes before wiping off.

Inactive ingredients

Water, Hydrochloric acid



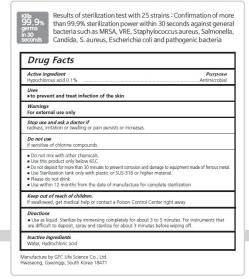
DAILY DOCTOR **Sterilization Spray Plus Up**

Kills 99.9% germs in 30 seconds

- $\cdot \ \, \text{For tableware, cutlery, cutting board, various kitchen utensils}$
- · For kindergarten, daycare center, newborn room, nursery
- · For bedding, table, chair, sofa, floor, wall, toilet, door handle
- · For hand washing sterilization to prevent pathogen infection

Ecofriendly sterilization disinfectant

16.9 fl.oz (500ml)





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DAILY DOCTOR HYPOCHLOROUS ACID DISINFECTANT LIQUID

hypochlorous acid spray

Product TypeHUMAN OTC DRUGItem Code (Source)NDC:76836-002

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength		
HYPO CHLO ROUS ACID (UNII: 712K4CDC10) (HYPO CHLO RITE ION - UNII: T5UM7HB19N)	HYPOCHLOROUS ACID	0.0001g in 1 mL		

Inactive Ingredients

inactive ingredients	
Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
HYDRO CHLO RIC ACID (UNII: QTT17582CB)	

Packaging

# Item Code	Package Description	Marketing Start Date	Marketing End Date	
1 NDC:76836-002- 01	60 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	05/14/2020		
2 NDC:76836-002- 02	500 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	05/14/2020		

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
unapproved drug other		05/14/2020		

Labeler - GFC Life Science Co., Ltd. (688325518)

Registrant - GFC Life Science Co., Ltd. (688325518)

Establishment				
Name	Address	ID/FEI	Business Operations	
GFC Life Science Co., Ltd.		688325518	manufacture(76836-002)	

Revised: 5/2020 GFC Life Science Co., Ltd.