DAILY DOCTOR STERILIZATION PLUS- sodium hypochlorite 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.

76836-001_Daily Doctor Sterilization Spray Plus

Active ingredient

Sodium Hypochlorite 0.2%

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



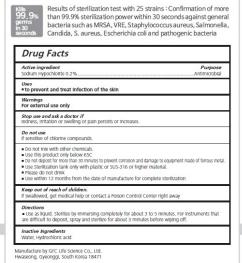
DAILY DOCTOR **Sterilization Spray Plus**

Kills 99.9% germs in 30 seconds

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

Ecofriendly sterilization disinfectant

16.9 fl.oz (500ml)







DAILY DOCTOR STERILIZATION PLUS

sodium hypochlorite spray

Product	Information
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Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:76836-001
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Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SODIUM HYPOCHLORITE (UNII: DY38 VHM5OD) (HYPOCHLORITE ION - UNII:T5UM7HB19N)	SODIUM HYPOCHLORITE	0.002 g in 1 mL

Inactive Ingredients		
Ingredient Name	Strength	
WATER (UNII: 059QF0KO0R)		
HYDRO CHLORIC ACID (UNII: QTT17582CB)		

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:76836-001- 01	60 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	04/29/2020	
2	NDC:76836-001- 02	500 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	04/29/2020	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		04/29/2020	

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(76836-001)		

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