

**DISINFECTANT LIQUID ANTIBACTERIAL- disinfectant liquid antibacterial spray spray**  
**Guangzhou Dedo Pharmaceutical 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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**Active Ingredient(s)**

HYPOCHLOROUS ACID 0.02%

**Purpose**

DISINFECTANT LIQUID

**Use**

SUITABLE FOR GENERAL OBJECT SURFACE,SKIN SURFACE,VEGETABLE AND FRUIT DISINFECTION ETC.EFFECTIVELY INHIBIT STAPHYLOCOCCUS AUREUS,CANDIDA ALBICANS AND ESCHERICHIA COLI.

**Warnings**

for external use only.

do not use in or near the eyes.In case of contact.rinse eyes thoroughly with water.Stop use and ask doctor if irritation or rash appears and lasts.

Keep out of reach for children.

If swallowed,get medical help or contact a Poison Control Center right away.

**Directions**

GENERAL OBJECT AND SKIN DISINFECTION:DIRECT SPRAYING OF PRODUCTS ON GENERAL OBJECTS OR SKIN SURFACES REQUIRING DISINFECTION;FRUITS AND VEGETABLES DISINFECTION:SPRAY DIRECTLY ON THE FRUIT AND VEGETABLE SURFACE AND FINSE WITH WATER AFTER 10 MINUTES

Children under 6 years of age should be supervised when using this product.

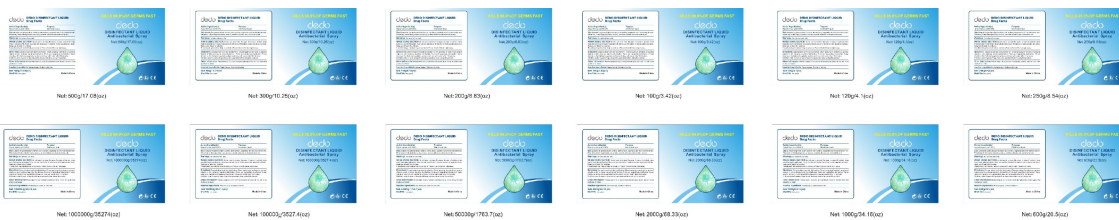
**Other information**

Please keep it in a cool and ventilated place.DO NOT MIX WITH OTHER PRODUCTS OR WATER.

**Inactive ingredients**

Water(Aqua),SODIUM CHLORIDE

**Package Label - Principal Display Panel**



## DISINFECTANT LIQUID ANTIBACTERIAL

disinfectant liquid antibacterial spray

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:74534-014
<b>Route of Administration</b>	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>HYPOCHLOROUS ACID</b> (UNII: 712K4CDC10) (HYPOCHLOROUS ACID - UNII:712K4CDC10)	HYPOCHLOROUS ACID	0.02 g in 100 g

### Inactive Ingredients

Ingredient Name	Strength
<b>SODIUM CHLORIDE</b> (UNII: 451W47IQ8X)	
<b>WATER</b> (UNII: 059QF0KO0R)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:74534-014-01	500 g in 1 BOTTLE; Type 0: Not a Combination Product	04/13/2020	
2	NDC:74534-014-12	600 g in 1 BOTTLE; Type 0: Not a Combination Product	04/13/2020	
3	NDC:74534-014-02	300 g in 1 BOTTLE; Type 0: Not a Combination Product	04/13/2020	
4	NDC:74534-014-03	200 g in 1 BOTTLE; Type 0: Not a Combination Product	04/13/2020	
5	NDC:74534-014-04	100 g in 1 BOTTLE; Type 0: Not a Combination Product	04/13/2020	
6	NDC:74534-014-05	120 g in 1 BOTTLE; Type 0: Not a Combination Product	04/13/2020	
7	NDC:74534-014-07	1000000 g in 1 BOTTLE; Type 0: Not a Combination Product	04/13/2020	
8	NDC:74534-014-06	250 g in 1 BOTTLE; Type 0: Not a Combination Product	04/13/2020	
9	NDC:74534-014-08	100000 g in 1 BOTTLE; Type 0: Not a Combination Product	04/13/2020	
10	NDC:74534-014-09	50000 g in 1 BOTTLE; Type 0: Not a Combination Product	04/13/2020	
11	NDC:74534-014-	2000 g in 1 BOTTLE; Type 0: Not a Combination Product	04/13/2020	

11	10	2000 g in 1 BOTTLE, Type 0: Not a Combination Product	04/13/2020	
12	NDC:74534-014-11	1000 g in 1 BOTTLE; Type 0: Not a Combination Product	04/13/2020	

<b>Marketing Information</b>			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		04/13/2020	

**Labeler** - Guangzhou Dedo Pharmaceutical Co.,Ltd (418394705)

**Registrant** - Guangzhou Dedo Pharmaceutical Co.,Ltd (418394705)

<b>Establishment</b>			
Name	Address	ID/FEI	Business Operations
Guangzhou Dedo Pharmaceutical Co.,Ltd		418394705	manufacture(74534-014)

Revised: 4/2020

Guangzhou Dedo Pharmaceutical Co.,Ltd