BACTERIA ZERO PREMIUM- peracetic acid liquid KUMSUNG E&C 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.

80138-204 BACTERIA ZERO PREMIUM

Active Ingredient[s]

Peracetic acid 0.015%

Purpose

Antiseptic

USE[s]

Fruits, Vegetables, Meat Sterilization(Food Additives), General Object surface Sterilization Sanitizer to help reduce bacteria that potentially can disease.

WARNINGS

For external use only. Flammable. Keep away form heat or flame Check the indications before use.

Do not inhale the contents that are sprayed or sprayed directly on a person.

When using this product

keep away from 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.

DIRECTIONS

- If the original solution is sprayed on a contaminated place, wipe it with a dry cloth or if it is naturally dried, harmful bacteria are immediately sterilized.
- Supervise children under 6 years of age when using this product to avoid swallowing.

OTHER INFORMATION

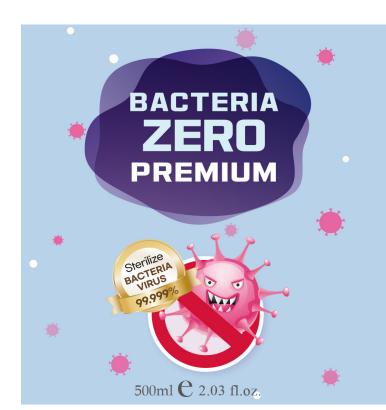
- Store between 1~30°C(33.8~86°F)
- Avoid freezing(-5°C) and excessive heat above 35°C(86°F)

INACTIVE INGREDIENTS

acetic acid, hydrogen peroxide, Phosphonic acid, purified water USP

Package Label





DRUG FACTS

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BACTERIA ZERO PREMIUM

peracetic acid liquid

Product Type HUMAN OTC DRUG Item Code (Source) NDC:80138-204

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Basis of Strength Ingredient Name Strength

PERACETIC ACID (UNII: 16KP12E1HD) (PERACETIC ACID - UNII:16KP12E1HD) PERACETIC ACID 0.015 g in 100 mL

Inactive Ingredients					
Ingredient Name	Strength				
PHOSPHOROUS ACID (UNII: 35V6A8JW8E)					
HYDROGEN PEROXIDE (UNII: BBX060AN9V)					
WATER (UNII: 059QF0KO0R)					
ACETIC ACID (UNII: Q40Q9N063P)					

P	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:80138- 204-01	60 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	01/01/2022		
٠,	NDC:80138-	500 mL in 1 BOTTLE, SPRAY; Type 0: Not a	01/01/2022		

204-02	Combination Product	01/01/2022			
Marketing Information					
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
unapproved drug		01/01/2022			
other		01/01/2022			

Labeler - KUMSUNG E&C CO., LTD (688082226)

Registrant - KUMSUNG E&C CO., LTD (688082226)

Establishment							
Name	Address	ID/FEI	Business Operations				
KUMSUNG E&C CO., LTD		688082226	manufacture(80138-204)				

Revised: 1/2022 KUMSUNG E&C CO., LTD