

AKTIVE MULTI PURPOSE ANTIBACTERIAL- hypochlorous acid liquid
Hall Global LLC

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.

Spray

This is a Antibacterilal Spray manufactured according to the Temporary Policy for Preparation of Certain Alcohol-Based Hand Sanitizer Products During the Public Health Emergency (CoViD-19); Guidance for Industry.

The Antibacterilal Spray is manufactured using only the following United States Pharmacopoeia (USP) grade ingredients in the preparation of the product (percentage in final product formulation) consistent with World Health Organization (WHO) recommendations:

- a. Hypochlorous Acid (EPA Reg# 96048-1) (USP or Food Chemical Codex (FCC) grade) (80%, volume/volume (v/v)) in an aqueous solution denatured according to Alcohol and Tobacco Tax and Trade Bureau regulations in 27 CFR part 20.
- b. Sterile distilled water or boiled cold water.
- c. Sodium Chloride

The firm does not add other active or inactive ingredients. Different or additional ingredients may impact the quality and potency of the product.

Active Ingredient(s)

Hypochlorous Acid (EPA Reg# 96048-1)

Purpose

Sanitizing/Antimicrobial Agent Solution is on the EPA N-List N-Product with Emerging Viral Pathogens and Human Coronavirus claims for use against SARS CoV-2

Use

Decreases bacteria on skin.

Warnings

For external use only.

Do not use

If you are allergic to the ingredient. Avoid contact to the eyes and skin. May irritate eyes or sensitive skin. If contact occurs rinse thoroughly with water.

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 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.

Stop use and ask a doctor if irritation or rash develops and continues for more than 72 hrs

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

away.

Directions

Adults and children 2 years and over: Spray on area and allow to dry without wiping. Children should be supervised when using and those under 2 years of age ask doctor before use. Discard empty bottle in trash receptacle.

Other information

- Store between 15-30C (59-86F)
- Avoid freezing and excessive heat above 40C (104F)

Inactive ingredients

purified water, Sodium Chloride

Package Label - Principal Display Panel

Multi-Purpose
Antibacterial
Spray

Anytime. Anywhere.
Stay Clean.

The primary ingredient hypochlorous acid (HOCl) is an all-natural compound produced by our body's immune system in our white blood cells, making it the most effective all-natural germ defender on the market without negative side effects. Multi-Purpose Antibacterial Spray contains NO harsh chemicals like sulfates, alcohol, and phosphates that can erode your skin health and cause your body to absorb toxic chemicals. The remaining two inactive ingredients contain sodium chloride (SALT) and oxygenated water, making this product all-natural and non-toxic.

AKTIVE™

Multi-Purpose Antibacterial Spray

Non-Irritating
pH Balanced
Non-Toxic
Non-Flammable
Non-Corrosive
Environmentally Friendly

No Formaldehyde
Biodegradable
Paraben Free
Alcohol Free
Fragrance Free
Pet-Friendly

KILLS
99%
OF GERMS



NDC 80546-001-15

8 FL OZ (226.8g)

000 mL NDC: 00000-000-00

AKTIVE MULTI PURPOSE ANTIBACTERIAL

hypochlorous acid liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:81156-002
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
HYPOCHLOROUS ACID (UNII: 712K4CDC10) (HYPOCHLOROUS ACID - UNII:712K4CDC10)	HYPOCHLOROUS ACID	0.25 mg in 100 mg

Inactive Ingredients

Ingredient Name	Strength
SODIUM CHLORIDE (UNII: 451W47IQ8X)	3.05 mg in 100 mg
WATER (UNII: 059QF0K00R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:81156-002-08	236 mg in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	03/30/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		03/30/2020	

Labeler - Hall Global LLC (117729725)**Registrant** - Hall Global LLC (117729725)**Establishment**

Name	Address	ID/FEI	Business Operations
Hall Global LLC		117729725	manufacture(81156-002)

Revised: 11/2020

Hall Global LLC