

**HAND SANITIZER- alcohol spray**  
**Axion Analytical Laboratories**

*Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.*

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**Sanity Hand Sanitizer**

This is a hand sanitizer 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 hand sanitizer 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. Alcohol (ethanol) (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. Glycerol (1.45% v/v).
- c. Hydrogen peroxide (0.125% v/v).
- d. Sterile distilled water or boiled cold water.

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)**

Alcohol 80% v/v. Purpose: Antiseptic

**Purpose**

Antiseptic, Hand Sanitizer

**Use**

Hand Sanitizer to help reduce bacteria that potentially can cause disease. For use when soap and water are not available.

**Warnings**

For external use only. Flammable. Keep away from heat or flame

**Do not use**

- in children less than 2 months of age
- on open skin wounds

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.

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

### Directions

- Place enough product on hands to cover all surfaces. Rub hands together until dry.
- Supervise children under 6 years of age when using this product to avoid swallowing.

### Other information

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

### Inactive ingredients

glycerin, hydrogen peroxide, purified water USP

### Package Label - Principal Display Panel

118 mL NDC: 75472-1234-4

Drug Facts	
<b>Active Ingredient</b>	<b>Purpose</b>
Ethyl Alcohol 80%.....	Antiseptic
<b>Uses</b>	
To decrease bacteria on the skin that could cause disease. Recommended for repeated use.	
<b>Warnings</b>	
For external use only. Flammable. Keep away from heat and flame.	
<b>When using this product:</b>	
Keep out of eyes. In case of contact with eyes, flush thoroughly with water. Avoid contact with broken skin. Do not inhale or ingest.	
<b>Stop use and ask a doctor if:</b> Irritation or redness develops or if condition persists more than 72 hours.	
Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.	
<b>Directions</b>	
Wet hands thoroughly with product and allow to dry without wiping. For children under 6, use only under adult supervision.	
<b>Other information:</b> Do not store above 105°Fahrenheit. May discolor some fabrics. Harmful to wood finishes and fabrics.	
<b>Inactive ingredients:</b> Purified water, Glycerin (plant-derived skin conditioner), Hydrogen Peroxide (anti-bacterial agent).	

**SANITY**

Trusted by hospitals,  
Loved by families.

**HAND SANITIZER  
POWER SPRAY**

kills >99.99% of illness-causing germs  
with the superior protection of  
80% ethyl alcohol

4 fl. oz. (118 mL)

Not tested on animals and no animal byproducts.  
Free of synthetic fragrances, parabens, triclosan,  
and benzethonium chloride.  
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www.sanity.company

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## HAND SANITIZER

alcohol spray

### Product Information

**Product Type**

HUMAN OTC DRUG

**Item Code (Source)**

NDC:75472-1234

Route of Administration TOPICAL

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	80 mL in 100 mL

### Inactive Ingredients

Ingredient Name	Strength
GLYCERIN (UNII: PDC6A3C0OX)	1.45 mL in 100 mL
HYDROGEN PEROXIDE (UNII: BBX060AN9V)	0.125 mL in 100 mL
WATER (UNII: 059QF0KO0R)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:75472-1234-4	118.29 mL 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
OTC monograph not final	part333A	03/15/2020	

**Labeler** - Axion Analytical Laboratories (016337854)

**Registrant** - Axion Analytical Laboratories (016337854)

### Establishment

Name	Address	ID/FEI	Business Operations
Axion Analytical Laboratories		016337854	analysis(75472-1234) , manufacture(75472-1234)

Revised: 4/2020

Axion Analytical Laboratories