

8 OZ HAND SANITIZER-LABS- alcohol spray
SunBeam Laboratories LLC

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.

8oz 237ml 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. 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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Directions

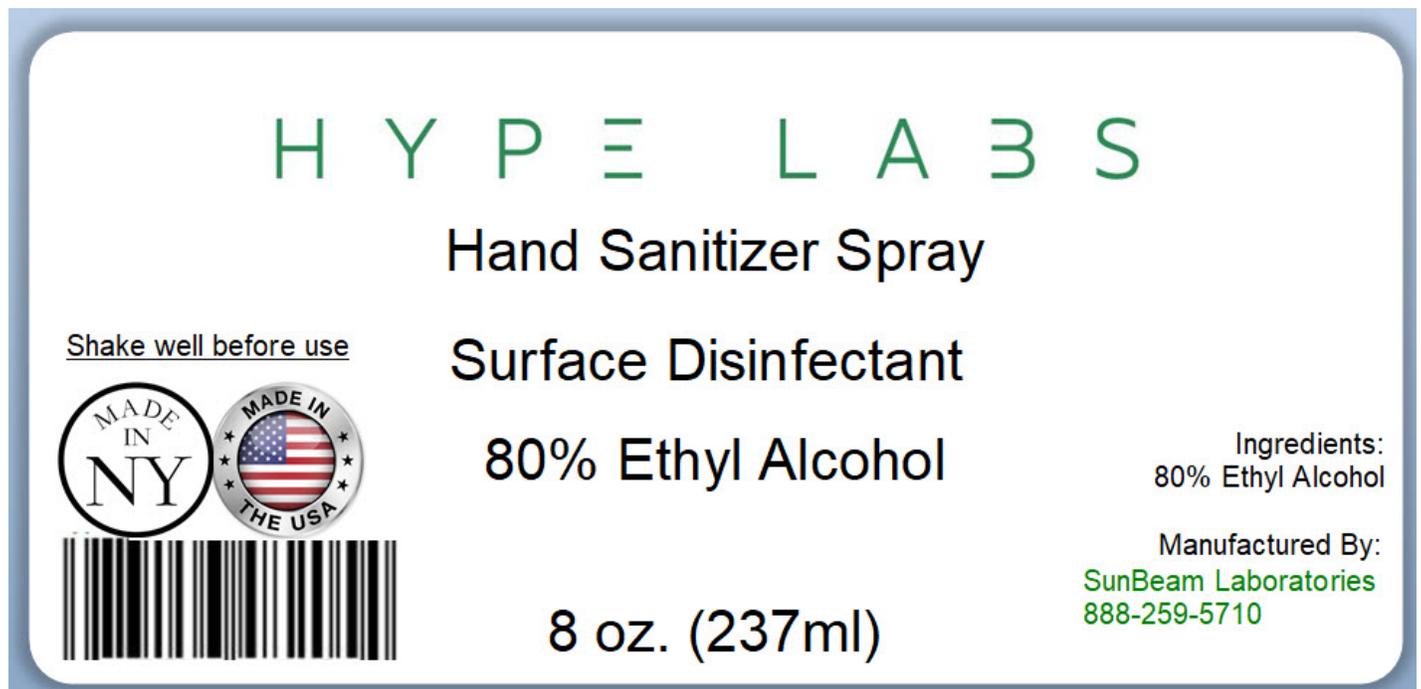
- Place enough product on hands to cover all surfaces. Rub hands together until dry.
- Supervise children under 6 years of age when using this product to avoid swallowing.

Other information

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

Inactive ingredients

purified water USP

Package Label - Principal Display Panel

The image shows the principal display panel of a product label. At the top, the brand name "HYPE LABS" is written in large, green, spaced-out capital letters. Below this, the product name "Hand Sanitizer Spray" is written in a black, sans-serif font. Underneath that, "Surface Disinfectant" is written in a similar black font. The active ingredient "80% Ethyl Alcohol" is prominently displayed in a large, bold, black font. At the bottom center, the volume "8 oz. (237ml)" is written in a black font. On the left side, there are two circular logos: one for "MADE IN NY" and another for "MADE IN THE USA" featuring an American flag. Below these logos is a standard 1D barcode. On the right side, the ingredients list "Ingredients: 80% Ethyl Alcohol" is shown in a small black font. Below that, the manufacturer information "Manufactured By: SunBeam Laboratories 888-259-5710" is written in a green font.

H Y P E L A B S

Hand Sanitizer Spray

Surface Disinfectant

80% Ethyl Alcohol

8 oz. (237ml)

Shake well before use

MADE IN NY

MADE IN THE USA

Ingredients:
80% Ethyl Alcohol

Manufactured By:
SunBeam Laboratories
888-259-5710

Drug Facts

Active ingredient[s]

Alcohol 80%v/v.....Antiseptic

Purpose

Use[s]

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Inactive ingredients purified water USP



STOP
CORONA



Hand Sanitizer Spray

Ingredients:
80% Ethyl Alcohol

Shake well before use

80% Denatured Ethyl Alcohol

Manufactured By:
SunBeam Laboratories



8 oz. (237ml)

888-259-5710

237 mL NDC: 75321-0008-2



Hand Sanitizer Spray
 Surface Disinfectant
 80% Ethyl Alcohol

Shake well before use



8 oz. (237ml)

Ingredients:
 80% Ethyl Alcohol

Manufactured By:
 SunBeam Laboratories
 888-259-5710

237 mL NDC: 75321-0008-3

8 OZ HAND SANI HYPE-LABS

alcohol spray

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:75321-0008
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	189 mL in 237 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0K00R)	47 mL in 237 mL

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:75321-0008-1	237 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	03/30/2020	
2	NDC:75321-0008-2	237 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	03/30/2020	
3	NDC:75321-0008-3	237 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	03/30/2020	

Labeler - SunBeam Laboratories LLC (105139335)

Establishment

Name	Address	ID/FEI	Business Operations
SunBeam Laboratories LLC		105139335	manufacture(75321-0008)

Revised: 4/2020

SunBeam Laboratories LLC