

WAVE HAND SANITIZER- alcohol liquid

Earthlite, 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.

wave 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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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

blood orange peel oil, broubon vanilla bean oil, cardamom seed extract, cedarwood, cinnamon bark oil, clove bud oil, glycerin, honey, hydrogen peroxide, jasmin, massoia bark extract, mimosa flower extract, patchouli leaf oil, purified water usp, sandalwood oil, sweet orange peel oil, ylang ylang flower oil

Package Label - Principal Display Panel

wave
HAND
SANITIZER
80% ALCOHOL
SOLUTION

- Topical Antiseptic
- CDC Recommended Formula
- FDA Registered
- Liquid Spray

8 oz (237 ml)

Drug Facts

Active ingredient[s]	Purpose
Alcohol 80% v/v	Antiseptic

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Inactive ingredients: Blood Orange Peel Oil, Bourbon Vanilla Bean Oil, Cardamom Seed Extract, Cedarwood, Cinnamon Bark Oil, Clove Bud Oil, Glycerin, Honey, Hydrogen peroxide, Jasmin, Massoia Bark Extract, Mimosa Flower Extract, Patchouli Leaf Oil, Purified water USP, Sandalwood Oil, Sweet Orange Peel Oil, Ylang Ylang Flower Oil.

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Made in the USA
Distributed by:
Tara Spa Therapy


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287 mL NDC: 77566-831-81

WAVE HAND SANITIZER

alcohol liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:77566-831
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:77566-831-81	287 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	07/29/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	07/29/2020	

Labeler - Earthlite, LLC (185340304)

Establishment

Name	Address	ID/FEI	Business Operations
Earthlite, LLC		185340304	manufacture(77566-831)

Revised: 7/2020

Earthlite, LLC