

HIGH PURITY HAND SANITIZER- alcohol gel

High Purity Natural Products 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.

Active Ingredient(s)

Alcohol 70% 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

DI Water, Vegetable Glycerin, Carbomer (ultrez 10), Diisopropylamine, Lemon Essential Oil

Package Label - Principal Display Panel

60 mL NDC: 75061-065-00

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NDC 75061-065-00 / Distributed by:
High Purity Natural Products, LLC / Southbridge, MA

HighPurityNaturaProducts.com



INSTANT HAND SANITIZER



The FDA has not approved for safety and efficacy. This product is not intended to diagnose, treat, cure, or prevent any disease.

DRUG FACTS

Active ingredient(s)	Purpose
Alcohol 70% w/v	Antiseptic

Use(s) Hand sanitizer to help reduce bacteria that potentially can cause disease.

For use when soap and water are not available.

Warnings

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Do not use:

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2 FL. OZ (60 ML)

HIGH PURITY HAND SANITIZER

alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:75061-065
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	70 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
GLYCERIN (UNII: PDC6A3C0OX)	0.5 mL in 100 mL
WATER (UNII: 059QF0K00R)	
DIISOPROPYLAMINE (UNII: BR9JLI40NO)	0.133 mL in 100 mL
LEMON OIL, DISTILLED (UNII: ET5GD00TRP)	0.15 mL in 100 mL
CARBOMER INTERPOLYMER TYPE A (5500 CPS) (UNII: 59TL3WG5CO)	0.274 mL in 100 mL

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:75061-065-00	60 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/25/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	05/31/2020	

Labeler - High Purity Natural Products LLC (114392832)

Establishment

Name	Address	ID/FEI	Business Operations
High Purity Natural Products LLC		114392832	manufacture(75061-065)

Revised: 7/2020

High Purity Natural Products LLC