

HAND SANITIZER- isopropyl alcohol liquid

Northshipco Inc.

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.

Hand Sanitizer IPA80% v/v

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. Isopropyl Alcohol (80%, 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)

Isopropyl 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.

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.

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



HAND SANITIZER

NON STERILE SOLUTION



Net Contents 128FL OZ (3.78 L)

MADE IN INDIA

DRUG FACTS

Active Ingredient

Isopropyl Alcohol 80% v/v

INACTIVE INGREDIENTS

Hydrogen Peroxide 0.125% (v/v) • Chlorhexidine Gluconate Soln IP 2.5% (v/v) • Glycerol 1.45% (v/v) • Aloe vera (Aloe Barbadensis Leaf) Extract 0.2% • Vitamin-E 0.02% • Phenoxyethanol 0.01% • Lemon Essential Oil 0.2% • Aqua Q.S. (Purified Water USP)

USES

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 fire or flame.

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.

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

OTHER INFORMATION

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

May discolor certain fabrics

Mfg. Lic. No.: NW(0300)/17/955

Batch No.:

Mfg. Date:

Best before 36 Months from the date of manufacturing

Not for Medical Use

Net Contents: 128FL OZ (3.78 L)



3.78L NDC: 80566-378-01

HAND SANITIZER

isopropyl alcohol liquid

Product Information

Product Type

HUMAN OTC DRUG

Item Code (Source)

NDC:80566-378

Route of Administration		TOPICAL		
Active Ingredient/Active Moiety				
Ingredient Name		Basis of Strength	Strength	
ISOPROPYL ALCOHOL (UNII: ND2M416302) (ISOPROPYL ALCOHOL - UNII:ND2M416302)		ISOPROPYL 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:80566-378-01	3780 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	09/15/2020	
Marketing Information				
Marketing Category		Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final		part333A	09/15/2020	

Labeler - Northshipco Inc. (029016807)

Establishment			
Name	Address	ID/FEI	Business Operations
Northshipco Inc.		029016807	manufacture(80566-378)

Revised: 9/2020

Northshipco Inc.