

CDC 70I- isopropyl alcohol gel
Friendo 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.

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. Isopropyl Alcohol (70%, v/v) in an aqueous solution denatured according to Alcohol and Tobacco Tax and Trade Bureau regulations in 27 CFR part 20.
- b. PVP (2.0% v/v).
- c. 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 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

PVP, purified water USP

Package Label - Principal Display Panel

| | |
|---|--|
|  | |
| Drug Facts | |
| Active Ingredient | Purpose |
| Isopropyl Alcohol (70%)..... | Antiseptic |
| Uses | |
| • For handwashing to decrease bacteria on the skin. • Recommended for repeated use. | |
| Warnings | |
| For external use only-hands Flammable. Keep away from heat and flame. When using this product • Keep out of eyes. In case of contact with eyes, flush thoroughly with water. • Avoid contact with broken skin. • Do not inhale or ingest. Stop use and ask a doctor if skin irritation develops. Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away. | |
| Directions • Wet hands thoroughly with product and allow to dry without wiping. • For children under 6, use only under adult supervision. • Not recommended for infants. | |
| Other Information • Do not store above 105° F. • May discolor some fabrics. • Harmful to wood finishes and plastics. | |
| Inactive Ingredients • 1-Ethenyl-2-pyrrolidone polymer, Water | |
| Distributed By: FRIENDO LLC 4285 TW King Road Southlake TX 76092 |  * L O T O O 1 0 5 * |

CDC 70I

isopropyl alcohol gel

Product Information

| | | | |
|-------------------------|----------------|--------------------|---------------|
| Product Type | HUMAN OTC DRUG | Item Code (Source) | NDC:74238-101 |
| Route of Administration | TOPICAL | | |

Active Ingredient/Active Moiety

| Ingredient Name | Basis of Strength | Strength |
|--|-------------------|--------------|
| ISOPROPYL ALCOHOL (UNII: ND2M416302) (ISOPROPYL ALCOHOL - UNII:ND2M416302) | ISOPROPYL ALCOHOL | 2.8 L in 4 L |

Inactive Ingredients

| Ingredient Name | | Strength | | |
|---------------------------------|--|--|----------------------|--------------------|
| POVIDONE K30 (UNII: U725QWY32X) | | 0.08 L in 4 L | | |
| WATER (UNII: 059QF0KO0R) | | 1.12 L in 4 L | | |
| Packaging | | | | |
| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
| 1 | NDC:74238-101-07 | 1041 L in 1 CONTAINER, FLEXIBLE INTERMEDIATE BULK; Type 0: Not a Combination Product | 03/31/2020 | |
| 2 | NDC:74238-101-08 | 1249 L in 1 CONTAINER, FLEXIBLE INTERMEDIATE BULK; Type 0: Not a Combination Product | 03/31/2020 | |
| Marketing Information | | | | |
| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date | |
| OTC monograph not final | part333A | 03/31/2020 | | |

Labeler - Friendo LLC (926569711)

Registrant - Friendo LLC (926569711)

| Establishment | | | |
|----------------------|---------|-----------|------------------------|
| Name | Address | ID/FEI | Business Operations |
| Friendo LLC | | 926569711 | manufacture(74238-101) |