**ISOPROPYL ALCOHOL 99 PERCENT- isopropyl alcohol liquid**
**All Pharma 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.*

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**Isopropyl Alcohol 99 Percent**

## Active Ingredient

Isopropyl Alcohol 99% by volume

## Purpose

First aid antiseptic

## Use

First aid to help prevent the risk of infection in.

- minor cuts
- scrapes
- burns

## Warnings

For external use only.

- Flammable, keep way from spark, heat and flame.

## Ask a doctor before use for

- deep wounds
- animal bites
- serious burns

## When using this product

• Do not inhale

• Do not apply over large areas of the body

• Do not use longer than 1 week

## Stop use and ask a doctor if

condition persists or gets worse

**Keep out of reach of children.**

In case of accidental ingestion, seek professional assistance or contact a Poison Control Center immediately.

**Directions**

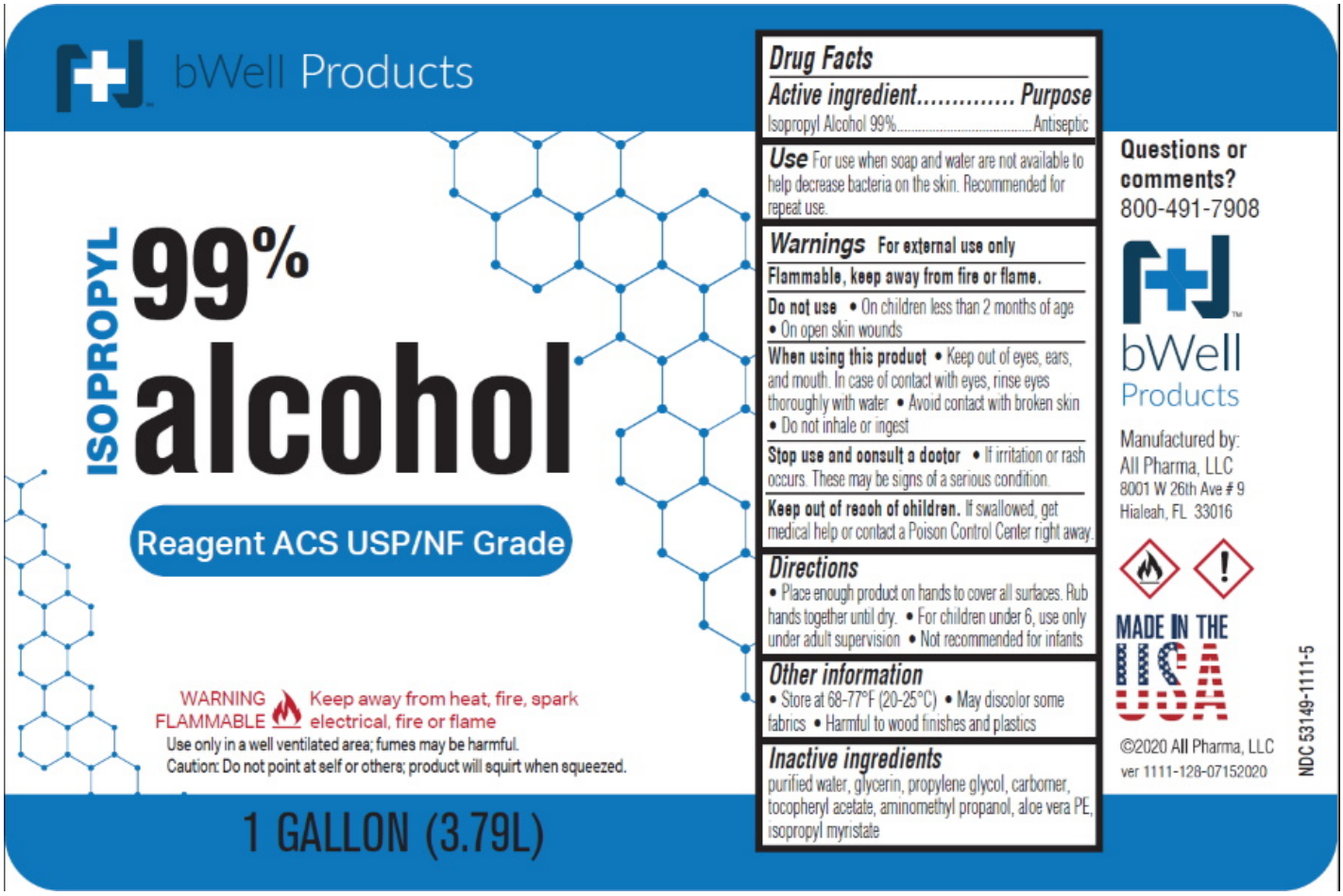
- clean the affected area.
- apply a small amount of this product on the affected area 1 to 3 times daily.
- may be covered with sterile bandage.
- if bandaged, let it dry first.

**Inactive Ingredient**

purified water

**Principal Display Panel**

**ISOPROPYL**
**RUBBING**
**ALCOHOL 99%**
**USP**
***First Aid Antiseptic***
**NDC 53149-1111-1 Bottle of 16 fl oz (1 pt) 473 mL**
**NDC 53149-1111-3 Bottle of 32 fl oz (1 qt) 946 mL**
**NDC 53149-1111-5 Bottle of 128 fl oz (1 Gallon) 3.79 L**




# ISOPROPYL ALCOHOL 99 PERCENT

isopropyl alcohol liquid

| Product Information | | | |
|---|---|---|---|
| **Product Type** | HUMAN OTC DRUG | **Item Code (Source)** | NDC:53149-1111 |
| **Route of Administration** | TOPICAL | | |

| Active Ingredient/Active Moiety | | |
|---|---|---|
| **Ingredient Name** | **Basis of Strength** | **Strength** |
| **ISOPROPYL ALCOHOL** (UNII: ND2M416302) (ISOPROPYL ALCOHOL - UNII:ND2M416302) | ISOPROPYL ALCOHOL | 990 mg in 100 mL |

| Inactive Ingredients | |
|---|---|
| **Ingredient Name** | **Strength** |
| **WATER** (UNII: 059QF0KO0R) | |

| Packaging | | | | |
|---|---|---|---|---|
| **#** | **Item Code** | **Package Description** | **Marketing Start Date** | **Marketing End Date** |

| | | | | |
|---|---|---|---|---|
| **1** | NDC:53149-1111-1 | 473 mL in 1 BOTTLE; Type 0: Not a Combination Product | 01/01/2020 | |
| **2** | NDC:53149-1111-3 | 946 mL in 1 BOTTLE; Type 0: Not a Combination Product | 01/01/2020 | |
| **3** | NDC:53149-1111-5 | 3790 mL in 1 BOTTLE; Type 0: Not a Combination Product | 01/01/2020 | |

## Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|---|---|---|---|
| OTC monograph not final | part333A | 01/01/2020 | |

## Labeler - All Pharma LLC (117605075)

## Registrant - All Pharma LLC (117605075)

## Establishment

| Name | Address | ID/FEI | Business Operations |
|---|---|---|---|
| All Pharma LLC | | 117605075 | manufacture(53149-1111) |

Revised: 1/2020 All Pharma LLC