

**ASPICARE- isopropyl alcohol solution**  
**American Private Label Products**

*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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**Active Ingredient**

Isopropyl Alcohol 70% v/v.

**Purpose**

Antiseptic

**Use**

Hand Sanitizer to help reduce bacteria on the skin

**Warnings**

**Flammable. Keep away from fire or flame. For external use only.**

**When using this product**

do not use in or near the eyes. In case of contact, rinse eyes thoroughly with water.

**Stop use and ask a doctor if**

irritation or rash appears or lasts

**Keep out of reach of children.**

If swallowed, get medical help or contact a Poison Control Center right away.

**Directions**

- Put enough product in your palm to cover hands and rub hands together briskly until dry
- Children under 6 years of age should be supervised when using ASPICARE Hand sanitizer

**Storage**

- Store below 110°F (43°C)
- May discolor certain fabrics or surfaces

**Inactive ingredients**

Water (Aqua), Acrylates Copolymer, Aloe Barbadensis Leaf Juice, Glycerin, Isosteareth-200 Linoleate, Disodium EDTA, Fragrance, Triethanolamine

**Other Information**

Hand sanitizers provide a convenient alternative when hand washing with plain soap and water is

unavailable. The Centers for Disease Control and Prevention advises that washing hands with plain soap and running water is one of the most important steps consumers can take to avoid getting sick and to prevent spreading infections to others. If soap and water are not available, the CDC recommends using an alcohol-based hand sanitizer. Hand sanitizers should not replace hand washing with soap and water. FDA conducts ongoing review of OTC antiseptic active ingredients to determine whether these ingredients are safe and effective for their intended uses.

**Principal Display Panel**

NDC 70735-071-01

AspiCare

HAND SANITIZER

KILLS 99% OF GERMS\*

Moisturizer Leaves Hands Smooth

MADE IN USA

1 FL OZ (30 mL)

NDC 70735-071-02

AspiCare

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MADE IN USA

2 FL OZ (60 mL)

NDC 70735-071-03

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3 FL OZ (89 mL)

NDC 70735-071-04

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MADE IN USA

4 FL OZ (118 mL)

NDC 70735-071-06

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MADE IN USA

6 FL OZ (177 mL)

NDC 70735-071-08

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MADE IN USA

8 FL OZ (236 mL)

NDC 70735-071-16

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MADE IN USA

16 FL OZ (473 mL)

NDC 70735-071-32

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32 FL OZ (946 mL)

## Drug Facts

Active ingredient	Purpose
Isopropyl Alcohol 70% v/v.....	Antiseptic

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Poison Control Center right away.

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Manufactured for:  
ASPICARE PRODUCTS  
Franklin NJ  
Made in the USA

www.aspicareproducts.com 8



# HAND SANITIZER

KILLS 99%  
OF GERMS\*

- Moisturizer Leaves  
Hands Smooth



8 FL OZ (236 mL)

NDC-XXXX-XXX-XX

## ASPICARE

isopropyl alcohol solution

### Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:70735-071
Route of Administration	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ISOPROPYL ALCOHOL (UNII: ND2M416302) (ISOPROPYL ALCOHOL - UNII:ND2M416302)	ISOPROPYL ALCOHOL	70 mL in 100 mL

## Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0K00R)	
BUTYL ACRYLATE/METHYL METHACRYLATE/METHACRYLIC ACID COPOLYMER (18000 MW) (UNII: JZ1374NL9E)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
GLYCERIN (UNII: PDC6A3C0OX)	
TROLAMINE (UNII: 9O3K93S3TK)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70735-071-01	30 mL in 1 BOTTLE; Type 0: Not a Combination Product	12/04/2020	
2	NDC:70735-071-02	60 mL in 1 BOTTLE; Type 0: Not a Combination Product	12/04/2020	
3	NDC:70735-071-03	89 mL in 1 BOTTLE; Type 0: Not a Combination Product	12/04/2020	
4	NDC:70735-071-04	118 mL in 1 BOTTLE; Type 0: Not a Combination Product	12/04/2020	
5	NDC:70735-071-06	177 mL in 1 BOTTLE; Type 0: Not a Combination Product	12/04/2020	
6	NDC:70735-071-08	236 mL in 1 BOTTLE; Type 0: Not a Combination Product	12/04/2020	
7	NDC:70735-071-16	473 mL in 1 BOTTLE; Type 0: Not a Combination Product	12/04/2020	
8	NDC:70735-071-32	946 mL in 1 BOTTLE; Type 0: Not a Combination Product	12/04/2020	

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	12/04/2020	

**Labeler** - American Private Label Products (051667474)

**Registrant** - American Private Label Products (051667474)

Revised: 12/2020

American Private Label Products