

**DOP ISOPROPYL RUBBING ALCOHOL 70%- isopropyl alcohol liquid  
Omega & Delta Co., Inc.**

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**DOP Isopropyl Rubbing Alcohol 70%**

***Drug Facts***

***Active ingredient***

Isopropyl alcohol 70%

***Purpose***

First aid antiseptic

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

**Use** helps prevent the risk of infection in minor cuts, scrapes and burns

***Warnings For external use only***

**Flammable keep away from fire or flame, heat, spark, electrical**

**Ask a doctor before use** for deep wounds, animal bites or serious burns

**When using this product** • do not get into eyes • do not apply over large area of the body • do not use longer than 1 week

**Stop use and ask a doctor if** the condition persists or gets worse

***Directions*** • clean the affected area • apply 1 to 3 times daily

***Inactive ingredient*** Purified water

***Other information*** • does not contain, nor is intended as a substitute for grain or ethyl alcohol • will produce serious gastric disturbances if taken internally

**Manufactured by:**

***Omega & Delta Co., Inc.***

**Carolina, P.R. 00984**

# DOP

# 70% Isopropyl Rubbing Alcohol

first aid antiseptic

- \* For Treatment of Minor Cuts and Scrapes
- \* First Aid Antiseptic
- \* Topical Sanitizer
- \* Body Rub / Massage

**Warning Flammable keep away from fire or flame**

16 FL OZ (1PT) 473 mL

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Distributed by:  
**Omega & Delta Co., Inc.**  
Carolina, P.R. 00984

## DOP ISOPROPYL RUBBING ALCOHOL 70%

isopropyl alcohol liquid

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:51048-005
<b>Route of Administration</b>	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: 059QF0KO0R)	

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:51048-005-16	473 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/01/2000	
2	NDC:51048-005-32	946 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/01/2000	
3	NDC:51048-005-04	118 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	01/21/2025	

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003	01/01/2000	

**Labeler** - Omega & Delta Co., Inc. (090317793)

## Establishment

Name	Address	ID/FEI	Business Operations
Omega & Delta Co., Inc.		090317793	manufacture(51048-005)

Revised: 1/2026

Omega & Delta Co., Inc.