**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 ISOPROPYL RUBBING ALCOHOL 70%

isopropyl alcohol liquid

| Product Information | | | |
|---|---|---|---|
| **Product Type** | HUMAN OTC DRUG | **Item Code (Source)** | NDC:51048-005 |
| **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: 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 | |

## 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: 11/2024 Omega & Delta Co., Inc.