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

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.

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	Marketing End

#	Item Code	Package Description	Date	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 not final	part333A	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: 11/2020 Omega & Delta Co., Inc.