

ISOPROPYL RUBBING ALCOHOL 70- isopropyl alcohol solution
Delon Laboratories (1990) Ltd.

Isopropyl Alcohol 70%

Active ingredient

Isopropyl alcohol 70% (by volume)

Purpose

First aid antiseptic

Uses

- first aid to help prevent the risk of infection in minor cuts, scrapes, and burns

Warnings

For external use only

- if taken internally serious gastric disturbances will result

Flammable, keep away from fire or flame

Ask a doctor before use if you have

- deep or puncture wounds, animal bites, or serious burns

When using this product

- do not get into eyes
- do not apply over large areas of the body
- do not use longer than 1 week unless directed by a doctor

Stop use and ask a doctor if

- the condition persists or gets worse

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

Directions

- clean the affected area
- may be covered with a sterile bandage
- if bandaged, let dry first

- apply a small amount of this product on the area 1 to 3 times daily

Other information

- store at room temperature
- does not contain, nor is intended as a substitute for grain or ethyl alcohol

Inactive ingredient

purified water

Drug Facts	
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NDC 61734-012-10



70%^{USP}

ISOPROPYL ALCOHOL



FIRST AID ANTISEPTIC

• First aid to help protect against infection in minor cuts, scrapes, and burns

16.9 fl. oz.





ASIN: B01UJKRN4 | 24050-1

Made in Canada.
Manufactured for:
RED Medical Supplies,
Vancouver, BC, Canada
V6H 1E3.
1-778-927-5647
redmedsupplies.com

SPACE FOR LOT # AND EXPIRY
DO NOT PRINT

ISOPROPYL RUBBING ALCOHOL 70

isopropyl alcohol solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:61734-012
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:61734-012-01	473 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/04/2014	02/07/2025
2	NDC:61734-012-02	118 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/04/2014	12/04/2014
3	NDC:61734-012-03	250 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/04/2014	12/04/2014
4	NDC:61734-012-04	1000 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/04/2014	12/04/2014
5	NDC:61734-012-05	3780 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/04/2014	12/04/2014
6	NDC:61734-012-06	100 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	10/04/2014	10/20/2022
7	NDC:61734-012-07	355 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/04/2014	12/04/2014
8	NDC:61734-012-08	946 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/04/2014	12/04/2014
9	NDC:61734-012-09	174 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	10/04/2014	10/31/2019
10	NDC:61734-012-10	500 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/27/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003	05/07/2010	

Labeler - Delon Laboratories (1990) Ltd. (248364184)

Establishment

Name	Address	ID/FEI	Business Operations
Laboratoires Delon		208896216	label(61734-012) , manufacture(61734-012) , pack(61734-012)

Revised: 10/2025

Delon Laboratories (1990) Ltd.