

ISOPROPYL ALCOHOL- isopropyl alcohol liquid
BROSANT GROUP LLC

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.

Active Ingredient

Isopropyl alcohol (70%conc)

Purpose

First aid antiseptic

Warnings

For external use only.

-if taken internally, serious gastric disturbances will result

Flammable keep away from fire or flamme

-use only in a well-ventilated area; fumes may be toxic

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 one week unless directed by a doctor

Stop use and ask a doctor if

condition persists or gets worse

keep out of reach of children

In case of ingestion, get medical help or contact a Poison Control Center right away (1-800-222-1222)

Directions

- clean affected area
- apply small amount of this product on the area 1-3 times daily
- if bandaged, let dry first
- may be covered with a sterile bandage

Other information

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

Inactive ingredient

purified water

Directions

- clean affected area
- apply small amount of this product on the area 1-3 times daily
- if bandaged, let dry first
- may be covered with a sterile bandage



Bioliving

70% isopropyl alcohol

warning flammable

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

use only in a well-ventilated area; fumes may be harmful

32FL OZ (1QT) 946 mL

ISOPROPYL ALCOHOL

isopropyl alcohol liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:83123-0019
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)	30 mL in 100 mL

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:83123-0019-1	946 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/17/2023	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	01/17/2023	

Labeler - BROSANT GROUP LLC (117037683)

Revised: 1/2023

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