

ALLURE ISOPROPYL RUBBING ALCOHOL WITH WINTERGREEN- isopropyl alcohol liquid

Universal Distribution Center LLC

ALLURE ISOPROPYL RUBBING ALCOHOL WITH WINTERGREEN

Active Ingredients

Isopropyl alcohol (50% conc.)

Purpose

Topical antimicrobial

Uses

- to decrease germs in minor cuts and scrapes.
- helps relieve minor muscular aches due to exertion.

Warnings

For external use only

- flammable, keep away from fire or flame, heat, spark, electrical.

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 body
- do not use longer than one week unless directed by a doctor

Stop using this product if

- condition persists or gets worse

Keep this and all drugs out of the reach of children

In case of accidental ingestion, seek professional assistance or contact a Poison control center (1-800-222-1212) immediately

Directions

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

Other information

- store at room temperature
- does not contain, nor is intended as a substitute for grain or ethyl alcohol, will produce serious gastric disturbance if taken internally

Inactive ingredient

purified water

Packaging

U

50% ISOPROPYL
with Wintergreen & Glycerin

Rubbing Alcohol

FIRST AID ANTISEPTIC

For Rubbing & Massaging

Use only in a well ventilated area;
Fumes may be harmful.

WARNING FLAMMABLE!
Keep away from fire or flame!

12 FL. OZ. (355 mL)

U

Isopropyl rubbing alcohol
50% by volume

Drug Facts

Active ingredients (by volume):	Purpose
Isopropyl alcohol (50% conc.)	first aid antiseptic

Uses ■ first aid to help 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 if you have
■ deep punctured 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 using this product if
■ condition persists or gets worse

Keep this and all drugs out of the reach of children.
In case of accidental ingestion, seek professional assistance or contact a Poison control center (1-800-222-1222) immediately.

Directions
■ clean effected area
■ apply small amount of this product on the area 1-3 times daily
■ May be covered with a sterile bandage
■ If bandaged, let dry first

Other information
■ store at room temperature
■ does not contain, nor is intended as a substitute for grain or ethyl alcohol. will produce serious gastric disturbances if taken internally.

Inactive Ingredient purified water

TAMPER EVIDENT: DO NOT USE IF THE UNDER CAP PRINTED SAFETY FOIL IS BROKEN OR MISSING.

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Made in Jordan

Distributed by: Universal Distribution Center
96 Distribution Boulevard • Edison, NJ 08817

ALLURE ISOPROPYL RUBBING ALCOHOL WITH WINTERGREEN

isopropyl alcohol liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:52000-002	
Route of Administration	TOPICAL			
Active Ingredient/Active Moiety				
Ingredient Name		Basis of Strength	Strength	
ISOPROPYL ALCOHOL (UNII: ND2M416302) (ISOPROPYL ALCOHOL - UNII:ND2M416302)		ISOPROPYL ALCOHOL	50 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:52000-002-01	118 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/30/2012	
2	NDC:52000-002-02	177 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/30/2012	
3	NDC:52000-002-03	237 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/30/2012	
4	NDC:52000-002-04	296 mL in 1 BOTTLE, PLASTIC; Type 1: Convenience Kit of Co-Package	06/30/2012	
5	NDC:52000-002-05	355 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/30/2012	
6	NDC:52000-002-06	414 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/30/2012	
7	NDC:52000-002-07	473 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/30/2012	
Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC Monograph Drug	M003	06/30/2012		

Labeler - Universal Distribution Center LLC (019180459)