ISOPROPYL RUBBING ALCOHOL 50% WITH WINTERGREEN- isopropyl alcohol liquid
Universal Distribution Center LLC

Isopropyl Rubbing Alcohol 50% With Wintergreen

Active Ingredients (by volume)

Isopropyl alcohol (50% conc.)

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; 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 (1-800-222-1212) 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

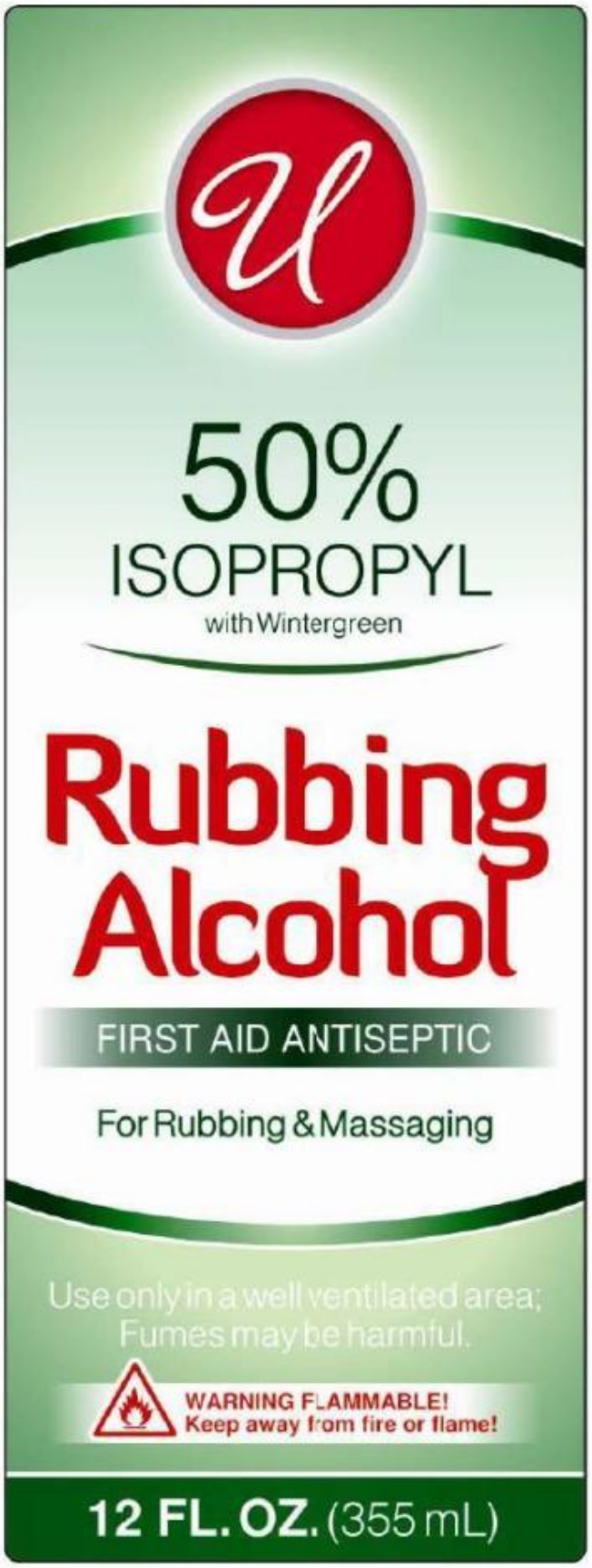
- store at room temperature
- does not contain, nor is intended as a substitute for grain or ethyl alcohol
- will produce serious gastric disturbance if taken internally

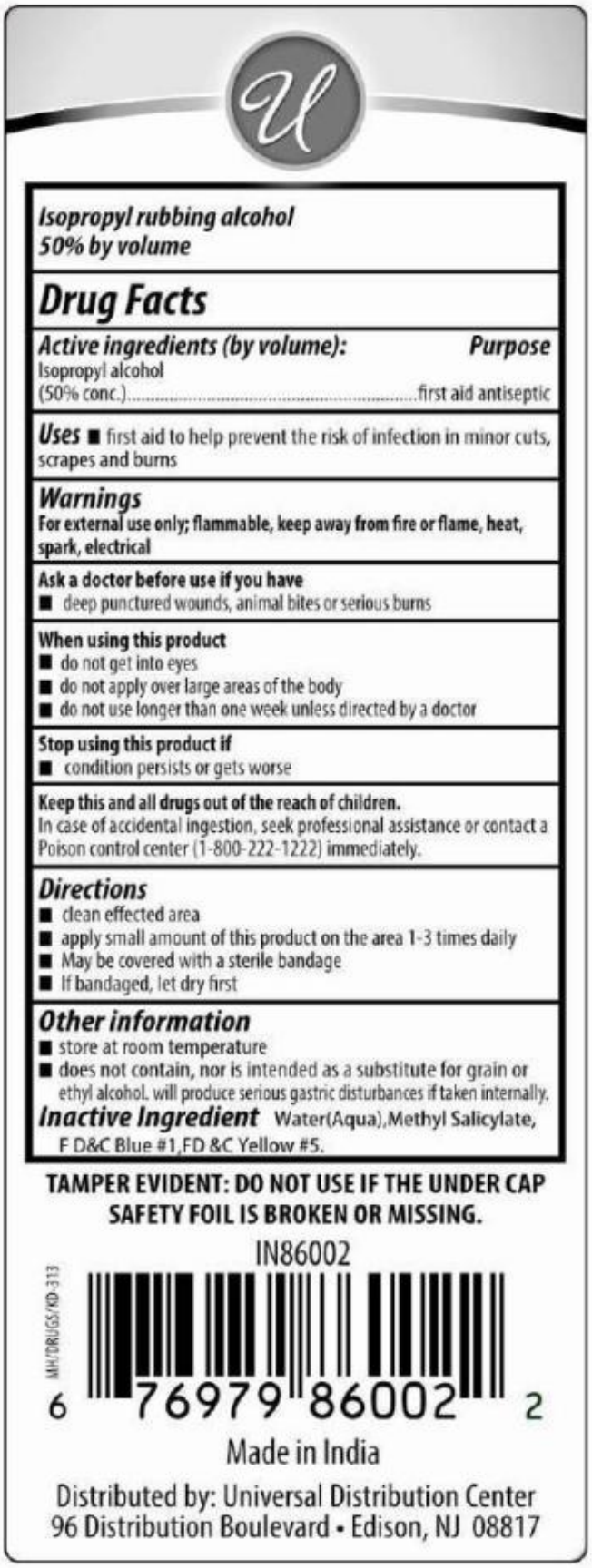
Inactive ingredient

Water(Aqua), Methyl Salicylate, FD&C Blue #1, FD&C Yellow #5

PRINCIPAL DISPLAY PANEL

ISOPROPYL RUBBING ALCOHOL 50% WITH WINTERGREEN
FIRST AID ANTISEPTIC
12 FL.OZ (355 mL)





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 Water(Aqua), Methyl Salicylate, F D&C Blue #1, FD &C Yellow #5.

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

IN86002

MH/DRUGS/KD-313

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Made in India

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

ISOPROPYL RUBBING ALCOHOL 50% WITH WINTERGREEN

isopropyl alcohol liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:52000-009
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)	
METHYL SALICYLATE (UNII: LAV5U5022Y)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:52000-009-01	118 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/15/2012	
2	NDC:52000-009-02	177 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/15/2012	
3	NDC:52000-009-03	237 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/15/2012	
4	NDC:52000-009-04	296 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/15/2012	
5	NDC:52000-009-05	355 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/15/2012	
6	NDC:52000-009-06	414 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/15/2012	
7	NDC:52000-009-07	473 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/15/2012	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003	03/15/2012	

Labeler - Universal Distribution Center LLC (019180459)

Registrant - Universal Distribution Center LLC (019180459)

Revised: 10/2024

Universal Distribution Center LLC