#### ISOPROPYL RUBBING ALCOHOL 70%- isopropyl alcohol liquid Universal Distribution Center 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.

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## Isopropyl Rubbing Alcohol 70%

#### Active Ingredients (by volume)

Isopropyl alcohol (70% 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

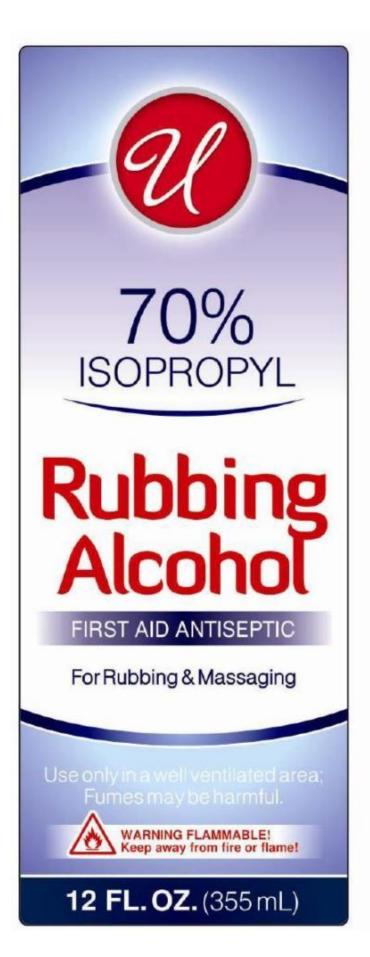
- store at room temperature
- does not contain, nor is intended as a substitute for grain or ethyl alcohol
- will produce serious gastric disturbance if taken internally

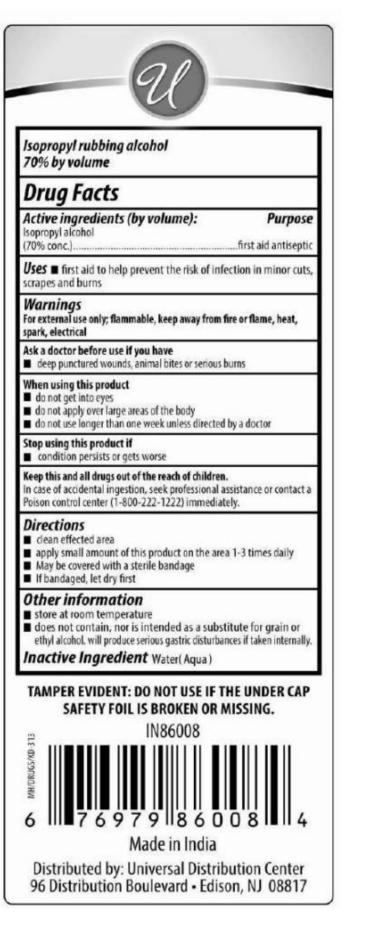
#### Inactive ingredient

Water(Aqua)

# PRINCIPAL DISPLAY PANEL

ISOPROPYL RUBBING ALCOHOL 70% FIRST AID ANTISEPTIC 12 FL.OZ (355 mL)





# **ISOPROPYL RUBBING ALCOHOL 70%**

FIUU	luct Info	ormation								
Product Type			HUMAN OTC DRUG	Item Code (Source)			NDC:52000-010			
Route of Administration			TOPICAL							
Active Ingredient/Active Moiety										
Ingredient Name					Basis of Strength			Strength		
ISOPROPYL ALCOHOL (UNII: ND2M416302) (ISOPROPYL ALCOHOL - UNII:ND2M416302)					ISOPROPYL ALCOHOL		7(	0 mL n 100 mL		
nact	tive Ingr	edients								
			redient Name			S	Streng	th		
VATE	<b>R</b> (UNII: 059	QF0KO0R)								
	aging	_		м	arketin	g Start	Mark	cetina End		
	aging m Code	Pa	ackage Description	м	arketin Dat	g Start te	Mark	ceting End Date		
# Ite	<b>cm Code</b>		TLE, PLASTIC; Type 0: Not	2			Mark			
#     Ite       1     NDC 010       2     NDC 010	em Code ::52000- -01 ::52000- -02	118 mL in 1 BO Combination Pro 177 mL in 1 BO Combination Pro	TTLE, PLASTIC; Type 0: Not oduct TTLE, PLASTIC; Type 0: Not oduct	a 03/ a 03/	Dat		Mark			
#     Ite       1     NDC 010       2     NDC 010       3     NDC 010	<b>c Code</b> <b>c:</b> 52000- -01 <b>c:</b> 52000- -02 <b>c:</b> 52000- -03	118 mL in 1 BOT Combination Pro 177 mL in 1 BOT Combination Pro 237 mL in 1 BOT Combination Pro	TTLE, PLASTIC; Type 0: Not oduct TTLE, PLASTIC; Type 0: Not oduct TTLE, PLASTIC; Type 0: Not oduct	a 03/ a 03/ a 03/ a 03/	<b>Dat</b> 15/2012		Mark			
#     Ite       1     NDC 010       2     NDC 010       3     NDC 010       4     NDC 010	em Code C:5200001 C:5200002 C:5200003 C:5200004	118 mL in 1 BO Combination Pro 177 mL in 1 BO Combination Pro 237 mL in 1 BO Combination Pro 296 mL in 1 BO Combination Pro	TTLE, PLASTIC; Type 0: Not oduct TTLE, PLASTIC; Type 0: Not oduct TTLE, PLASTIC; Type 0: Not oduct TTLE, PLASTIC; Type 0: Not oduct	a 03/ a 03/ a 03/ a 03/ a 03/	Dat 15/2012 15/2012		Mark			
#     Ite       1     NDC 010       2     NDC 010       3     NDC 010       4     NDC 010       5     NDC 010	em Code 5:5200001 5:5200002 5:5200003 5:5200004 5:5200005	118 mL in 1 BOT Combination Pro 177 mL in 1 BOT Combination Pro 237 mL in 1 BOT Combination Pro 296 mL in 1 BOT Combination Pro 355 mL in 1 BOT Combination Pro	TTLE, PLASTIC; Type 0: Not oduct TTLE, PLASTIC; Type 0: Not oduct TTLE, PLASTIC; Type 0: Not oduct TTLE, PLASTIC; Type 0: Not oduct TTLE, PLASTIC; Type 0: Not oduct	a 03/ a 03/ a 03/ a 03/ a 03/ a 03/	Dat 15/2012 15/2012 15/2012		Mark			
#     Ite       1     NDC 010       2     NDC 010       3     NDC 010       4     NDC 010       5     NDC 010	em Code C:5200001 C:5200002 C:5200003 C:5200004 C:5200005 C:5200005 C:5200006	118 mL in 1 BOT Combination Pro 177 mL in 1 BOT Combination Pro 237 mL in 1 BOT Combination Pro 296 mL in 1 BOT Combination Pro 355 mL in 1 BOT Combination Pro 414 mL in 1 BOT	TTLE, PLASTIC; Type 0: Not oduct TTLE, PLASTIC; Type 0: Not oduct TTLE, PLASTIC; Type 0: Not oduct TTLE, PLASTIC; Type 0: Not oduct TTLE, PLASTIC; Type 0: Not oduct	a 03/	Dat 15/2012 15/2012 15/2012 15/2012		Mark			
#     Ite       1     NDC 010       2     NDC 010       3     NDC 010       4     NDC 010       5     NDC 010       6     NDC 010	em Code C:5200001 C:5200002 C:5200003 C:5200004 C:5200005 C:5200006 C:5200006 C:5200006	118 mL in 1 BOT Combination Pro 177 mL in 1 BOT Combination Pro 237 mL in 1 BOT Combination Pro 296 mL in 1 BOT Combination Pro 355 mL in 1 BOT Combination Pro 414 mL in 1 BOT	TTLE, PLASTIC; Type 0: Not oduct TTLE, PLASTIC; Type 0: Not	a   03/     a   03/	Dat 15/2012 15/2012 15/2012 15/2012 15/2012		Mark	ceting Enc Date		
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Labeler - Universal Distribution Center LLC (019180459)

**Registrant -** Universal Distribution Center LLC (019180459)

Establishment									
Name	Address	ID/FEI	<b>Business Operations</b>						
Anicare Pharmaceuticals Pvt. Ltd		916837425	manufacture(52000-010)						

Revised: 11/2022

Universal Distribution Center LLC