# DONE 70 ETHYL ALCOHOL- alcohol liquid Olein Recovery Corporation

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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# Done 70% Ethyl Alcohol

# **Drug Facts**

# **Active Ingredient (by volume)**

Ethyl Alcohol 70%

# **Purpose**

first aid antiseptic

### Uses

• first aid to help prevent the risk of infection in minor cut, scrapes and burns

# **Warnings**

# For external use only

- flammable, keep away from fire or flame
- if taken internally serious gastric disturbances will result

# Ask a doctor before use if you have

• deep or punture 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 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 immediately.

# 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**

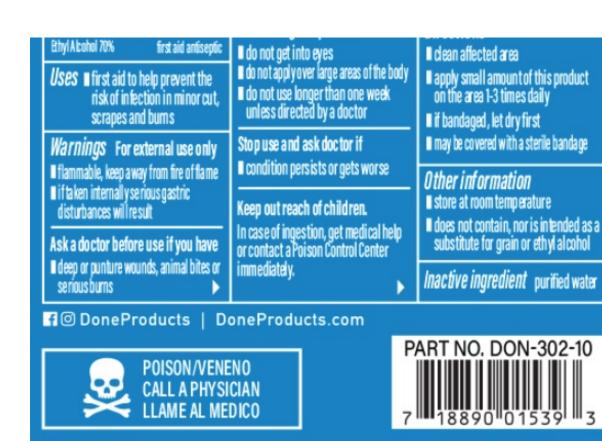
purified water

Package Labeling: 236.5ml



Package Labeling: 295.7ml

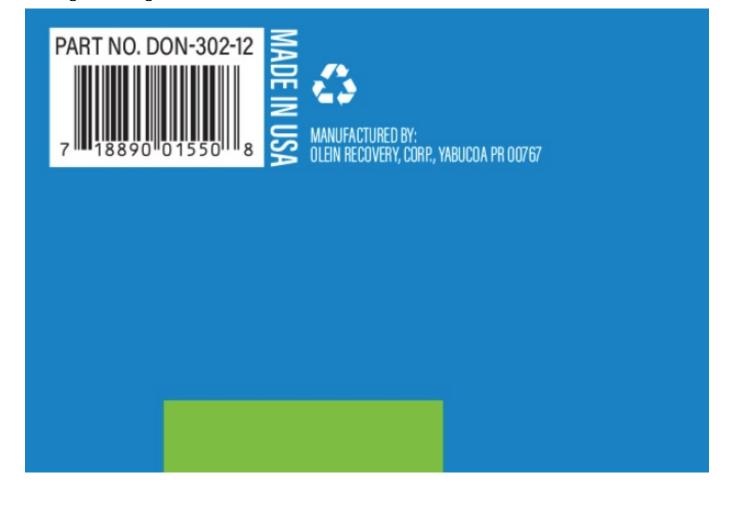




Package Labeling: 354.8ml

**MADEIN USA** 

10 fl. oz. (295.7 mL)



MANUFACTURED BY:

OLEIN RECOVERY, CORP., YABUCOA PR 00767

MAJJE IN USA 12 fl. oz. (354.8 mL)

# First Aid Antiseptic



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# Drug Facts (continued)

# Directions

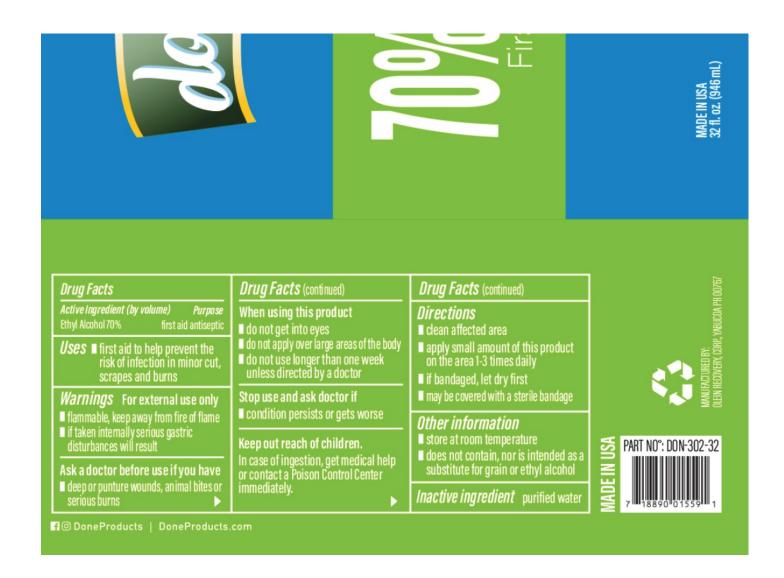
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Other information

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Package Labeling: 946ml





Package Labeling: 3785.4ml



First Aid Antiseptic

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# **Drug Facts** (continued)

■ clean affected area

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# Other information

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Inactive ingredient purified water



# MADE IN USA



128 fl oz (3785.4 mL) 🚓

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# **DONE 70 ETHYL ALCOHOL**

alcohol liquid

### **Product Information**

HUMAN OTC DRUG NDC:77142-010 Product Type Item Code (Source)

**Route of Administration** TOPICAL

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	0.7 mL in 1 mL		

Inactive Ingredients					
Ingredient Name	Strength				
WATER (UNII: 059QF0KO0R)					

P	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:77142-010-01	236.5 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/15/2020		
2	NDC:77142-010- 02	295.7 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/15/2020		
3	NDC:77142-010- 03	354.8 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/15/2020		
4	NDC:77142-010- 04	946 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/15/2020		
5	NDC:77142-010- 05	3785.4 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/15/2020		

Marketing Information					
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
OTC monograph not final	part333E	05/15/2020			

# Labeler - Olein Recovery Corporation (188543446)

Revised: 5/2020 Olein Recovery Corporation