ETHYL RUBBING ALCOHOL 70 PERCENT- ethyl alcohol liquid
Humco Holding Group, Inc.

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.

Private Label Ethyl Rubbing Alcohol 70 Percent USP

Drug Facts

Active Ingredient

Ethyl Alcohol 70% by volume

Purpose

First aid antiseptic

Use

First aid to help prevent the risk of infection in.

- minor cuts
- scrapes
- burns

Warnings

For external use only.

- Flammable, keep way from spark, heat and flame.

Ask a doctor before use for

- deep wounds
- animal bites
- 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 1 week

Stop use and ask a doctor if

condition persists or gets worse

Keep out of reach of children.

In case of accidental ingestion, seek professional assistance or contact a Poison Control Center immediately.

Directions

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

Other Information

- store at room temperature
- will produce serious gastric disturbances if taken internally.

Inactive Ingredient

Acetone 5%

Methyl isobutyl ketone 1%

bitrex

purified water

Principal Display Panel

NDC 46122-329-43
RUBBING ALCOHOL
ETHYL ALCOHOL 70%
USP
First Aid Antiseptic
16 fl oz (1 pt) 473 mL

ETHYL RUBBING ALCOHOL 70 PERCENT

ethyl alcohol liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0395-9100
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	700 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
ACETONE (UNII: 1364PS73AF)	
METHYL ISOBUTYL KETONE (UNII: U5T7B88CNP)	
DENATONIUM BENZOATE (UNII: 4YK5Z54AT2)	
WATER (UNII: 059QF0KO0R)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:0395-9100-16	473 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/01/2016	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	01/01/2016	

Labeler - Humco Holding Group, Inc. (825672884)

Registrant - Humco Holding Group, Inc. (825672884)

Establishment

Name	Address	ID/FEI	Business Operations
Humco Holding Group, Inc.		825672884	manufacture(0395-9100), analysis(0395-9100), pack(0395-9100), label(0395-9100)

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