ISOPROPYL ALCOHOL- isopropyl alcohol liquid
Phoenix Pharmaceutical Inc./ Clipper Distributing, Inc.

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

Isopropyl Alcohol 99%

NDC 57319-431-09
Isopropyl Alcohol 99%
ANTISEPTIC, DISINFECTANT RUBEFACIENT

FOR ANIMAL USE ONLY

KEEP OUT OF REACH OF CHILDREN

Net Contents:

3.785 L (1 GAL)

PHOENIX

PHARMACEUTICAL, INC.

Manufactured for:

Clipper Distributing Company, LLC

St. Joseph, MO 64507

INDICATIONS:

For external use only as an antiseptic, disinfectant, and rubefacient.

TO MAKE A STANDARD SOLUTION (70%):

Dilute by adding 1 part water to 2 parts of this 99% Isopropyl Alcohol.

FOR EXTERNAL USE ONLY

DIRECTIONS:

Use full strength as topical antiseptic or for disinfections of instruments.

May be used for temporary relief of minor muscular aches or pain due to overexertion and fatigue. Apply full strength to affected area and massage briskly to stimulate circulation.

CAUTION:

In case of deep or puncture wounds or serious burns, consult a veterinarian. If redness, irritation, or swelling persists or increases, discontinue use and consult a veterinarian. Do not apply to irritated skin or if excessive irritation develops.

INGREDIENTS:

Isopropyl Alcohol ..... 99% w/v

Water ..... q.s.

WARNING:

If taken internally, serious gastric disturbance will result.

First Aid:

Induce vomiting or use stomach pump. Call a physician immediately. Avoid getting into eyes or on mucous membranes. In case of eye contact, flush thoroughly with water. Call a physician immediately.

STORE AT CONTROLLED ROOM TEMPERATURE BETWEEN 15°-30°C (59°-86°F).

KEEP CONTAINER CLOSED WHEN NOT IN USE.

WARNING! FLAMMABLE! KEEP AWAY FROM HEAT AND OPEN FLAME.

TAKE TIME OBSERVE LABEL DIRECTIONS

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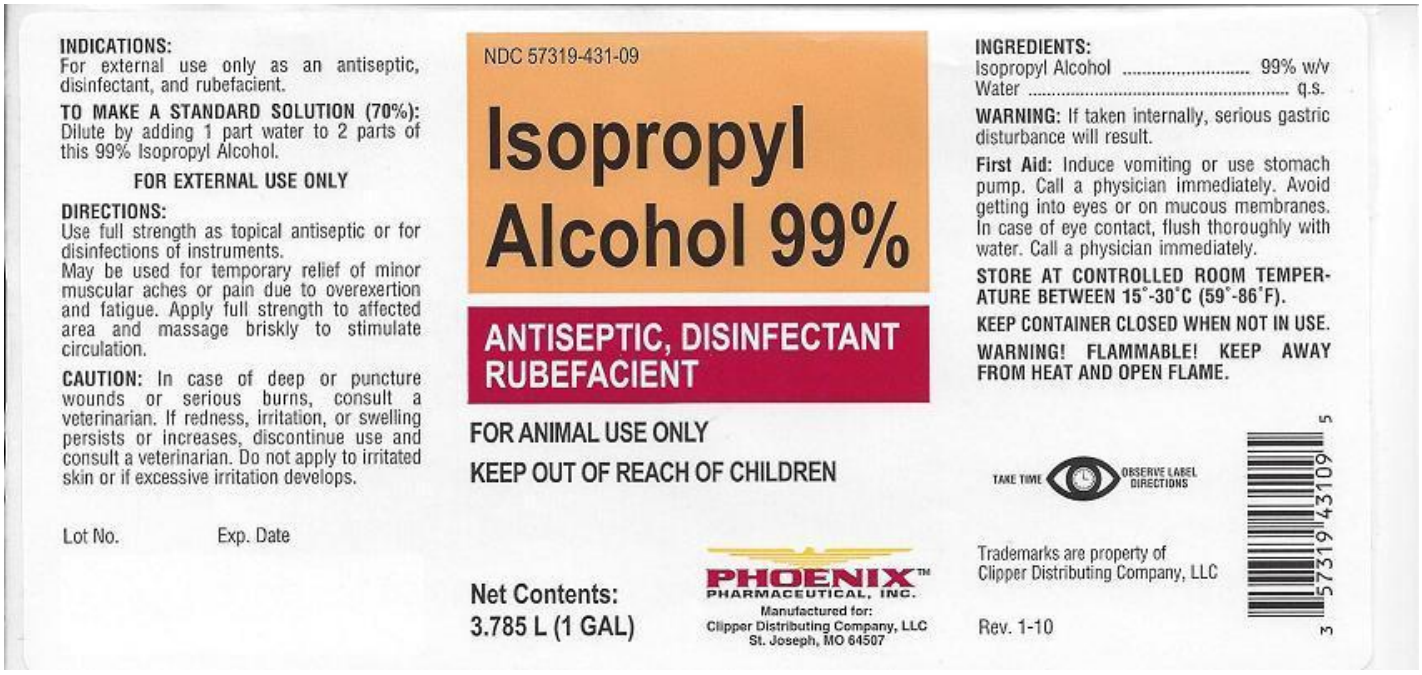
Rev 1-10



ISOPROPYL ALCOHOL

isopropyl alcohol liquid

Product Information			
Product Type	OTC ANIMAL DRUG	**Item Code (Source)**	NDC:57319-431
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**

ISOPROPYL ALCOHOL (UNII: ND2M416302) (ISOPROPYL ALCOHOL - UNII:ND2M416302)	ISOPROPYL ALCOHOL	99 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:57319-431-22	946 mL in 1 BOTTLE		
2	NDC:57319-431-09	3785 mL in 1 BOTTLE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
Unapproved drug other		11/15/2002	

Labeler - Phoenix Pharmaceutical Inc./ Clipper Distributing, Inc. (150711039)

Revised: 9/2013 Phoenix Pharmaceutical Inc./ Clipper Distributing, Inc.