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 70%

NDC 57319-430-09
Isopropyl Alcohol 70%
FOR EXTERNAL USE ONLY
KEEP OUT OF REACH OF CHILDREN

NET CONTENTS:

3.785L (1 GALLON)

PHOENIX™

PHARMACEUTICAL, INC.

Manufactured for:

Clipper Distributing Company, LLC

St. Joseph, MO 64507

For Veterinary Use Only

Scrub hands and arms with soap and water, rinse with water, then scrub with alcohol solution for disinfecting.

External Solution for use as a topical antiseptic. may also be used for temporary relief of minor muscular aches or pain due to overexertion and fatigue. Apply full strength directly to affected area, wet thoroughly and massage briskly to stimulate circulation.

Store at controlled room temperature between 15° and 30°C (59°-86°F).

Manufactured by:
Ameri-Pac Inc.
St. Joseph, MO 64502

Rev. 10-09

TAKE TIME OBSERVE LABEL DIRECTIONS

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Clipper Distributing Company, LLC

ACTIVE INGREDIENT:

Isopropyl Alcohol 70%

INERT INGREDIENT

Deionized Water 30%
Total 100%

WARNING

For external use only. If taken internally, serious gastric disturbance will result. Avoid contact with eyes. In case of eye contact, flush thoroughly with water. Call a physician.

First Aid:

Induce vomiting or use stomach pump.

CAUTION

FLAMMABLE
KEEP AWAY FROM FIRE OR FLAME

KEEP CONTAINER CLOSED WHEN NOT IN USE

ISOPROPYL ALCOHOL

isopropyl alcohol liquid

Product Information			
Product Type	OTC ANIMAL DRUG	**Item Code (Source)**	NDC:57319-430
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ISOPROPYL ALCOHOL (UNII: ND2M416302) (ISOPROPYL ALCOHOL - UNII:ND2M416302)	ISOPROPYL ALCOHOL	70 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:57319-430-22	946 mL in 1 BOTTLE		
2	NDC:57319-430-09	3785 mL in 1 BOTTLE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
Unapproved drug other		01/29/2002	

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

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