

ISOPROPYL ALCOHOL- isopropyl alcohol liquid

Durvet, 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.

Durvet 70% Alcohol Solution

NDC 30798-786-35

durvet

70% ALCOHOL SOLUTION

Contains 70% Isopropyl Alcohol

Rubefacient

www.durvet.com

NET CONTENTS: ONE GALLON (3.785 L)

FOR ANIMAL USE ONLY

INDICATIONS: Helps relieve minor muscular aches due to overexertion.

DIRECTIONS: Apply liberally and rub in.

CAUTION:

Harmful if swallowed.

WARNING:

FOR EXTERNAL USE ONLY. Keep out of reach of children. If taken internally, severe gastric disturbances will result. In case of accidental ingestion, call a physician or poison control center immediately. Causes eye irritation. In case of contact, immediately flush eyes with water and call a physician. May cause skin irritation. Avoid contact with eyes, skin, mucous membranes and clothing. Wash hands thoroughly after handling.

TAKE TIME OBSERVE LABEL DIRECTIONS

Manufactured for:

DURVET, INC.

Blue Springs, Missouri 64014

Iss. 07-09

KEEP OUT OF REACH OF CHILDREN

CONTENTS:

Isopropyl Alcohol 70%

Purified Water 30%

This preparation is made from Isopropyl Alcohol and does not contain, nor is it sold as a substitute for, ethyl or grain alcohol.

STORAGE:

Store at controlled room temperature between 15° - 30°C (59°-86°F). Keep tightly closed, protect from light.

WARNING! FLAMMABLE!

KEEP AWAY FROM HEAT AND OPEN FLAME

Lot No. Exp. Date

Label

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ISOPROPYL ALCOHOL

isopropyl alcohol liquid

Product Information

Product Type	OTC ANIMAL DRUG	Item Code (Source)	NDC:30798-786
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:30798-786-35	3785 mL in 1 BOTTLE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
Unapproved drug other		08/10/2009	

Labeler - Durvet, Inc. (056387798)

Revised: 9/2013

Durvet, Inc.