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.

99% ALCOHOL SOLUTION

Contains 99% Isopropyl Alcohol

Rubefacient

FOR ANIMAL USE ONLY

KEEP OUT OF REACH OF CHILDREN

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 sever 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.

WARNING! FLAMMABLE! KEEP AWAY FROM HEAT AND OPEN FLAME

ACTIVE INGREDIENTS
Isopropyl Alcohol 99%

INACTIVE INGREDIENTS
Purified Water 1%

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^{o}-30^{o}C (59^{o}-86^{o}F). Keep tightly closed, protect from sunlight.

ISOPROPYL ALCOHOL

isopropyl alcohol liquid

Product Information			
Product Type	OTC ANIMAL DRUG	**Item Code (Source)**	NDC:30798-467
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 L in 100 L

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	1 L in 100 L

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:30798-467-35	3.785 L in 1 JUG		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		01/01/2010	

Labeler - Durvet, Inc (056387798)

Revised: 9/2014 Durvet, Inc