

MAGNALAX- laxative powder, for solution
Vedco, 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.

MAGNALAX POWDER

MILD LAXATIVE/ANTACID

FOR ANIMAL USE ONLY

KEEP OUT OF REACH OF CHILDREN

INDICATIONS

A mild laxative, antacid for use in ruminant animals with simple indigestion.

DIRECTIONS

Mature bovine, stir one pound of Magnalax Powder into one gallon of water and mix thoroughly.

Administer as a drench or via stomach tube. For sheep, goats and calves administer one pint to one quart depending on size of animals. Frequent or continued use should be avoided.

WARNING

Milk that has been taken from animals during treatment and for 12 hours (1 milking) after the latest treatment must not be used for food.

CONTENTS

Each pound contains:

Magnesium Hydroxide 10.35 oz. in a flavored base.

One pound contains magnesium hydroxide equivalent to 1 gallon milk of magnesia.

DO NOT STORE ABOVE 30° C (86°F)

TAKE TIME OBSERVE LABEL DIRECTIONS

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M-0737-09
VINV-MAGP-25LB

Rev. 06-22

Distributed By
VEDCO, INC.
St. Joseph, MO 64507



ACTIVE INGREDIENTS:

Each pound contains:
Magnesium Hydroxide 10.35 oz.

INACTIVE INGREDIENTS: in a flavored base.

One pound contains magnesium hydroxide equivalent to 1 gallon milk of magnesia.

Do Not Store Above 30°C (86°F).



Lot No. Exp. Date



MAGNALAX

laxative powder, for solution

Product Information

Product Type	OTC ANIMAL DRUG	Item Code (Source)	NDC:50989-061
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MAGNESIUM HYDROXIDE (UNII: NBZ3QY004S) (MAGNESIUM CATION - UNII:T6V3LHY838)	MAGNESIUM HYDROXIDE	293.7 g in 454 g

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:50989-061-26	454 g in 1 JAR		
2	NDC:50989-061-40	11350 g in 1 PAIL		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		03/18/1998	

