

RENAPLUS- potassium gluconate powder
MWI/VetOne

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.

RenaPlus™ Powder

□Potassium Gluconate

Keep out of reach of children.□

□CAUTION: □ Federal law restricts this drug to use by or on the order of a licensed veterinarian.

□FOR USE IN ANIMALS ONLY

□Each 0.65 g (1/4 level teaspoon) contains a minimum:

Potassium gluconate.....2 mEq (468 mg) in a palatable base.

Dosage:

The suggested dose of RenaPlus for adult cats and dogs is 0.65 g (1/4 level teaspoon) per 10 lb (4.5 kg) body weight twice daily with food. Dosage may be adjusted to satisfy patient's need.

□Precautions:

Use with caution in the presence of cardiac disease, particularly in digitalized patients or in the presence of renal disease.

Lot No.:

Exp. Date:

Warning:

Do not administer in diseases where high potassium levels may be encountered, such as severe renal insufficiency or adrenal insufficiency.

□Indication:

For use as a supplement in potassium deficient states in cats and dogs.

Storage:

□Store tightly closed container in a dry place at a temperature not exceeding 35°C (95°F).

Distributed by: MWI

Boise, ID 83705

(888) 694-8381

PRINCIPAL DISPLAY PANEL - 4 oz Bottle

NDC: 13985-591-51

RenaPlus™

Potassium Gluconate

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VETone®

V 502060

Net Contents: 4 oz (113 g)

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www.VetOne.net
Rev. 07/13

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Lot No.:

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TAKE TIME



OBSERVE LABEL DIRECTIONS

RENAPLUS

potassium gluconate powder

Product Information

Product Type	PRESCRIPTION ANIMAL DRUG	Item Code (Source)	NDC:13985-591
Route of Administration	Oral		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
POTASSIUM GLUCONATE (UNII: 12H3K5QKN9) (POTASSIUM CATION - UNII:295O53K152)	POTASSIUM GLUCONATE	468 mg in 0.65 g

Inactive Ingredients

Ingredient Name	Strength
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SILICON DIOXIDE (UNII: ETJ7Z6XBU4)

Product Characteristics

Color	brown	Score	
Shape		Size	
Flavor	LIVER	Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:13985-591-51	113 g in 1 BOTTLE, PLASTIC		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		06/06/2014	

Labeler - MWI/VetOne (019926120)

Establishment

Name	Address	ID/FEI	Business Operations
Neogen Corporation-Mercer		042125879	analysis, manufacture, label

Revised: 11/2014

MWI/VetOne