

RENACARE K- potassium gluconate gel
Cosco International, 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](#).

PHP Animal Health RenaCare K

Product Facts

Potassium Gluconate2 mEq (468 mg) per 2.5g

INACTIVE INGREDIENTS:

Biotin, Citric Acid, Cyanocobalamin, Folic Acid, Maple Flavor, Niacinamide, Pantothenic Acid, Potassium Sorbate, Purified Water, Pyridoxine Hydrochloride, Riboflavin, Sodium Benzoate, Sodium Carboxymethylcellulose, Sucralose, Thiamine Hydrochloride.

CAUTIONS:

If animal's condition worsens or does not improve, stop product administration and consult your veterinarian.

Safe use in pregnant animals or animals intended for breeding has not been proven.

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

Do not administer to cats or dogs with acute, oliguric, anuric or very advanced renal failure.

Do not administer in other diseases where high potassium levels may be encountered, such as adrenal insufficiency, acute dehydration or urethral obstruction.

INTENDED USE:

FOR USE IN ADULT CATS & DOGS ONLY

Recommended for Renal Health Support

DIRECTIONS FOR USE:

Administer orally: Adult dogs and cats give 1/2 teaspoon per 10 lbs. (4.5 kg) body weight, twice daily.

Adjust amount based on veterinary recommendations.

WARNINGS:

FOR ANIMAL USE ONLY. KEEP OUT OF THE REACH OF CHILDREN AND ANIMALS. IN CASE OF ACCIDENTAL OVERDOSE, CONTACT A HEALTH PROFESSIONAL IMMEDIATELY. STORE AT A CONTROLLED ROOM TEMPERATURE.

QUESTIONS:

Made in U.S.A

Distributed by:

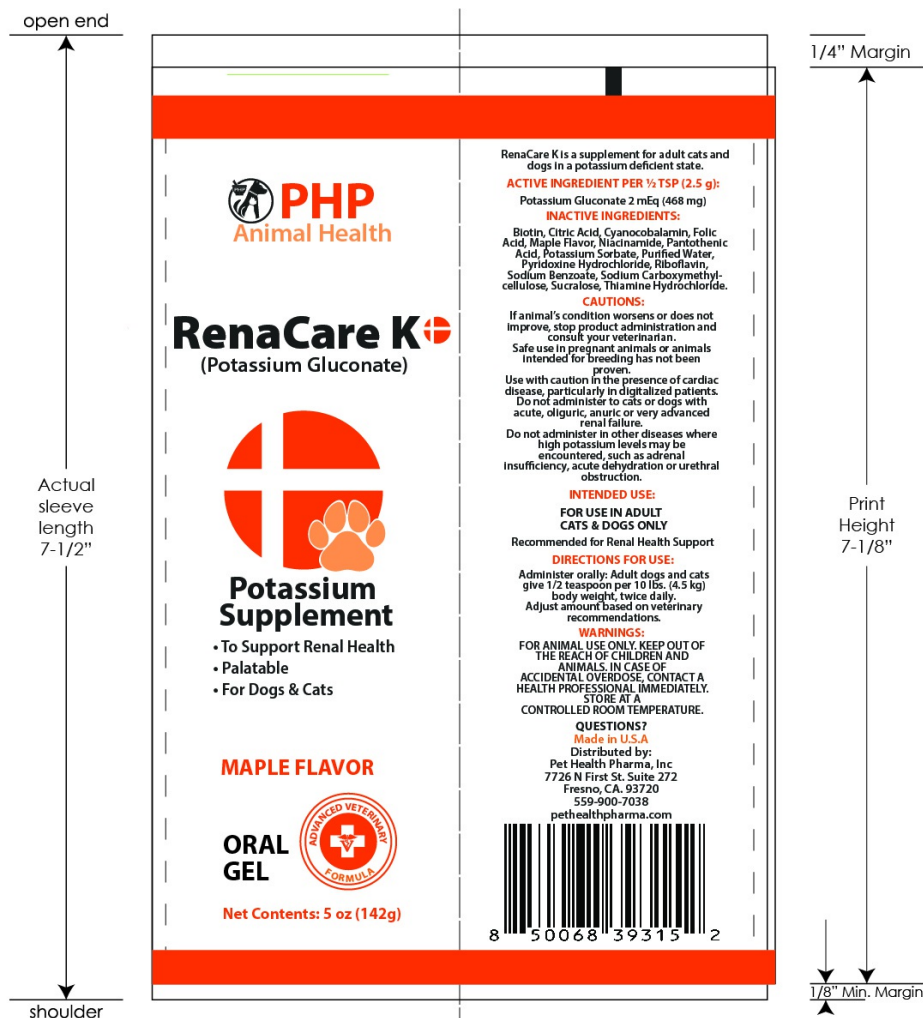
Pet Health Pharma, Inc.

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RENACARE K

potassium gluconate gel

Product Information

Product Type	OTC ANIMAL DRUG	Item Code (Source)	NDC:52261-7001
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
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Potassium Gluconate (UNII: 12H3K5QKN9) (POTASSIUM CATION - UNII:295O53K152)			Potassium Gluconate	468 mg in 1 g
Inactive Ingredients				
Ingredient Name				Strength
BIOTIN (UNII: 6SO6U10H04)				
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)				
CYANOCOBALAMIN (UNII: P6YC3EG204)				
FOLIC ACID (UNII: 935E97BOY8)				
NIACINAMIDE (UNII: 25X51I8RD4)				
PANTOTHENIC ACID (UNII: 19F5HK2737)				
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)				
WATER (UNII: 059QF0KO0R)				
PYRIDOXINE HYDROCHLORIDE (UNII: 68Y4CF58BV)				
RIBOFLAVIN (UNII: TLM2976OFR)				
SODIUM BENZOATE (UNII: OJ245FE5EU)				
CARBOXYMETHYLCELLULOSE SODIUM (UNII: K679OBS311)				
SUCRALOSE (UNII: 96K6UQ3ZD4)				
THIAMINE HYDROCHLORIDE (UNII: M572600E5P)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:52261-7001-1	142 g in 1 TUBE		
Marketing Information				
Marketing Category	Application Number or Monograph Citation		Marketing Start Date	Marketing End Date
unapproved drug other			05/01/2024	

Labeler - Cosco International, Inc. (016433141)

Registrant - Cosco International, Inc. (016433141)

Establishment			
Name	Address	ID/FEI	Business Operations
Cosco International, Inc.		016433141	manufacture, api manufacture