

CAL-DEX CMPK- cmpk injection, solution

Covetrus

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](#).

Cal-dex CMPK

COMPOSITION:

Each 500 mL of sterile aqueous solution contains.

Calcium (as calcium borogluconate, equivalent to calcium gluconate 23.2%)	10.8 g
Potassium (as potassium chloride)	8.0 g
Phosphorus (as sodium hypophosphite • H ₂ O)	2.5 g
Magnesium (as magnesium borogluconate)	1.6 g
Dextrose • H ₂ O	75.0 g

MILLIEQUIVALENTS PER LITER

Cations

Calcium	1,080 mEq/L
Potassium	410 mEq/L
Magnesium	261 mEq/L
Sodium	161 mEq/L

Anions

Borogluconate	1,341 mEq/L
Chloride	410 mEq/L
Hypophosphite	161 mEq/L

INDICATIONS:

For use as an aid in the treatment of hypocalcemia (parturient paresis, milk fever), hypomagnesemia (grass tetany), and other conditions associated with calcium, magnesium, phosphorus and potassium deficiencies in cattle.

CONTRAINDICATIONS:

Do not administer this product to animals showing signs of cardiac distress.

PRECAUTIONS:

Administration should be made slowly and with care to avoid adverse effects such as heart block or shock. Perivascular or subcutaneous deposition of hypertonic solutions may result in severe inflammation at the injection site.

CAUTION:

This product contains no preservatives. Use entire contents when first opened. Discard any unused solution.

DOSAGE AND ADMINISTRATION:

Warm solution to body temperature. The usual intravenous dose in cattle is 500 mL per 800 to 1,000 pounds of body weight.

Store between 15°C - 30°C (59°F - 86°F).

TAKE TIME OBSERVE LABEL DIRECTIONS

Keep Out of Reach of Children

For Animal Use Only

CAUTION:

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

NDC: 11695-6412-5

For treatment of milk fever in cattle

Volume: 16.907 fl oz (500 mL)

Questions? (855) 724-3461

Reorder #069177

18-805

RMS 92-1177

AH-069177-01

REV: 0719

Lot No.

Exp. Date

Distributed by:

Covetrus North America

400 Metro Place North

Dublin, OH 43017

covetrus.com

Manufactured by:

Nova-Tech, Inc.

Grand Island, NE 68801

Assembled in USA

Composición: Cada 500 mL de solución acuosa estéril contienen:

Calcio	10.8 g
(como borogluconato de calcio, equivalente a gluconato de calcio, 23.2%)	
Potasio (como cloruro de potasio)	8.0 g
Fósforo (como hipofosfito de sodio • H ₂ O)	2.5 g
Magnesio (como borogluconato de magnesio)	1.6 g
Dextrosa • H ₂ O	75.0 g

Milieuivalentes por litro

Cationes		Aniones	
Calcio	1,080 mEq/L	Borogluconato	1,341 mEq/L
Potasio	410 mEq/L	Cloruro	410 mEq/L
Magnesio	261 mEq/L	Hipofosfito	161 mEq/L
Sodio	161 mEq/L		

Indicaciones: Como ayuda en el tratamiento de hipocalcemia (paresis parturienta, fiebre de leche), hipomagnesemia (tétano de los pastos) y otras condiciones asociadas con deficiencias de calcio, magnesio, fósforo y potasio en los bovinos.

Contraindicaciones: No administrar este producto a animales que muestren señales de malestares cardíacos.

Precauciones: La administración debe realizarse lentamente y con cuidado afín de evitar efectos adversos como shock o bloqueo cardíaco. La deposición perivascular o subcutánea de soluciones hipertónicas puede resultar en inflamación severa en el lugar de la inyección.

Precaución: Este producto no contiene agentes conservantes. Usar todo el contenido después de abrir. Descartar cualquier solución no utilizada.

Dosis y administración: Elevar la temperatura de la solución a la del cuerpo. La dosis intravenosa usual para los bovinos es de 500 mL por cada 800 a 1000 libras EE.UU. (365 a 456 kg) de peso corporal. Almacénese entre 15° y 30° C (59° y 86° F).

Tomarse el tiempo requerido para observar las direcciones de la etiqueta.

№ para nuevo pedido 069177



NDC: 11695-6412-5

Cal-dex CMPK injection

 Cal-dex CMPK inyección

For treatment of milk fever in cattle
Para el tratamiento de la fiebre de leche en los bovinos

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Keep out of reach of children

Solamente para uso animal
Manténgase fuera del alcance de los niños

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Product Information

Product Type	PRESCRIPTION ANIMAL DRUG	Item Code (Source)	NDC:11695-6412
Route of Administration	INTRAVENOUS		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
Dextrose monohydrate (UNII: LX22YL083G) (ANHYDROUS DEXTROSE - UNII:5SL0G7R0OK)	Dextrose monohydrate	75.0 g in 500 mL

Potassium Chloride (UNII: 660YQ98I10) (POTASSIUM CATION - UNII:295O53K152)	POTASSIUM CATION	8.0 g in 500 mL
Calcium Gluconate Monohydrate (UNII: CZN0M5R31) (CALCIUM CATION - UNII:2M83C4R6ZB)	Calcium Gluconate Monohydrate	10.8 g in 500 mL
Phosphorus (UNII: 27YLU75U4W) (PHOSPHORUS - UNII:27YLU75U4W)	Phosphorus	2.5 g in 500 mL
Magnesium Gluconate (UNII: T42NAD2KHC) (MAGNESIUM CATION - UNII:T6V3LHY838)	Magnesium Gluconate	1.6 g in 500 mL

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11695-6412-5	500 mL in 1 BOTTLE, PLASTIC		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		03/17/2020	

Labeler - Covetrus (603750329)

Registrant - Nova-Tech, Inc. (196078976)

Establishment

Name	Address	ID/FEI	Business Operations
Nova-Tech, Inc.		196078976	manufacture, api manufacture

Revised: 3/2020

Covetrus