

CMPK- cmpk injection, solution
Nova-Tech, 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](#).

CMPK Injection

Each 500 mL Contains

Dextrose • H ₂ O.....	75.0 g
Calcium	10.8 g
(as calcium borogluconate)	
Potassium	8.0 g
(as potassium chloride)	
Phosphorus	2.5 g
(as sodium hypophosphite • H ₂ O)	
Magnesium	1.6 g
(as magnesium borogluconate)	

INDICATIONS

For use in cattle as an aid in treatment of conditions associated with calcium, magnesium, phosphorus, potassium or glucose deficiencies.

CONTRAINDICATIONS

DO NOT administer this product to animals showing signs of cardiac distress. Monitor animal's condition closely.

CAUTION

Entire contents should be used when first entered as this product contains no preservatives. Do not use if solution is not clear.
Discard any unused portion.

DOSAGE & ADMINISTRATION:

Cattle500 mL/800-1000 lbs body weight.

For intravenous administration only. Solution should be warmed to body temperature and administered slowly using aseptic precautions.

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.

Manufactured by:

Nova-Tech, Inc.

Grand Island, NE 68801 USA

18-805

RMS 92-336

NDC: 65207-805-50

Nova-Tech® Animal Health

Calcium-Magnesium-Phosphorus-Potassium-Dextrose

Steril Solution

Net Contents:

500 mL (16.91 fl oz)

Assembled in USA

Lot No.

Exp. Date

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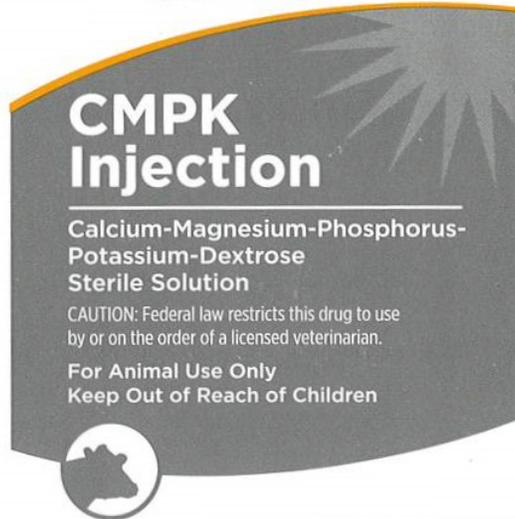
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Nova-Tech, Inc.
Grand Island, NE 68801 USA

18-805
RMS 92-336

NDC# 65207-805-50



CMPK Injection

Calcium-Magnesium-Phosphorus-Potassium-Dextrose Sterile Solution

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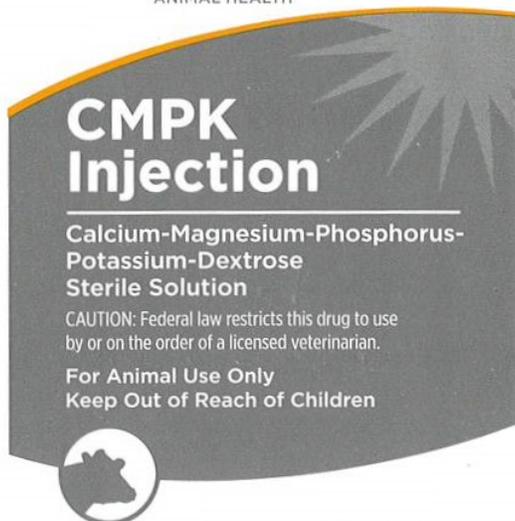
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Grand Island, NE 68801 USA

18-805
RMS 92-336

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CMPK

cmpk injection, solution

Product Information

Product Type	PREScription ANIMAL DRUG	Item Code (Source)	NDC:65207-805
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: 660 YQ98I10) (POTASSIUM CATION - UNII:295O53K152)	POTASSIUM CATION	8.0 g in 500 mL
Calcium Gluconate Monohydrate (UNII: CZN0MI5R31) (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:65207-805-50	500 mL in 1 BOTTLE, PLASTIC		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		03/18/2020	

Labeler - Nova-Tech, Inc. (196078976)

Registrant - Nova-Tech, Inc. (196078976)

Establishment

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

Revised: 3/2020

Nova-Tech, Inc.