

CALCIUM GLUCONATE- calcium gluconate 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.

CALCIUM GLUCONATE 23%

INDICATIONS

For use in cattle as an aid in treating uncomplicated milk fever (parturient paresis).

DOSAGE & ADMINISTRATION

For intravenous administration only.

Cattle 250-500 mL

Dosage may be repeated at 8 to 12 hour intervals if required.

TAKE TIME OBSERVE LABEL DIRECTIONS

Manufactured by:

Nova-Tech Inc.

Grand Island, NE 68801 USA

18-802

RMS 92-326

NDC# 65207-802-50

Nova-Tech[®] Animal Health

Calcium Gluconate 23%

Sterile Solution

Net Contents:

500 mL (16.91 fl oz)

Assembled in USA

Lot No.

Exp. Date

Each 500 mL Contains

Calcium Gluconate 23% w/v

(provides 10.71 g of Calcium* per 500 mL)

Water for Injection q.s.

*Present as salts or boryl esters of gluconic acid.

Store between 15°C-30°C (59°F-86°F).

FOR ANIMAL USE ONLY

KEEP OUT OF REACH OF CHILDREN

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Caution

Solution should be warmed to body temperature and administered slowly. Entire contents should be used upon entering, as this product contains no preservative. Discard any unused portion. Do not use if solution is cloudy or contains precipitate.

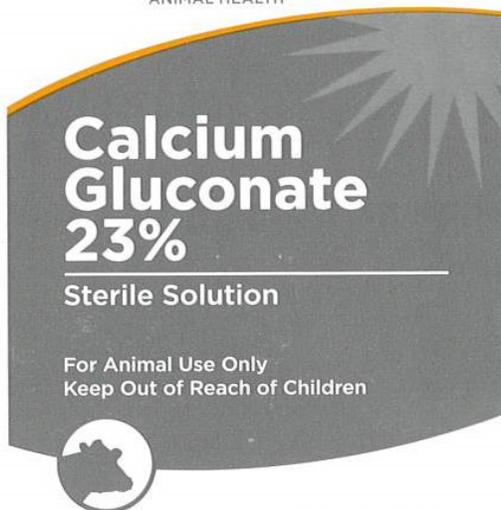
Note

Monitor animal's condition closely. Re-treatment may be necessary. If there is no noticeable improvement within 24 hours following treatment, consult your veterinarian. Aseptic precautions should be observed.

Manufactured by:

Nova-Tech, Inc. 18-802
Grand Island, NE 68801 USA RMS 92-326

NDC# 65207-802-50



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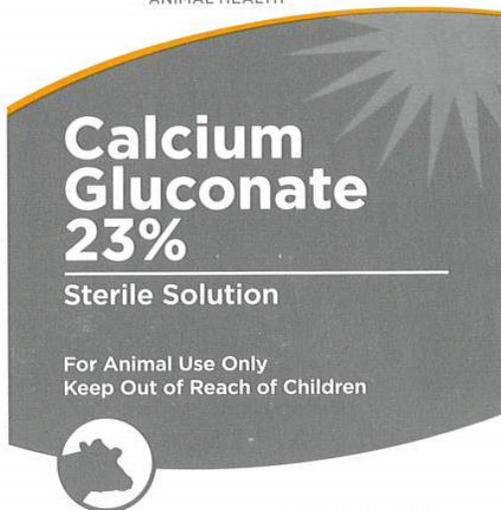
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CALCIUM GLUCONATE

calcium gluconate injection, solution

Product Information

Product Type	OTC ANIMAL DRUG	Item Code (Source)	NDC:65207-802
Route of Administration	INTRAVENOUS		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CALCIUM GLUCONATE (UNII: SQE6VB453K) (CALCIUM CATION - UNII:2M83C4R6ZB)	CALCIUM GLUCONATE	23 g in 100 mL

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:65207-802-50	500 mL in 1 BOTTLE, PLASTIC		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		03/19/2020	

Labeler - Nova-Tech, Inc. (196078976)**Registrant** - Nova-Tech, Inc. (196078976)**Establishment**

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

Revised: 3/2020

Nova-Tech, Inc.