Calcitat N 25 pro inf.

Authorised

- Calcium gluconate monohydrate
- Magnesium chloride hexahydrate
- Phosphorylcolamine
- Calcium hydroxide
- Calcium borogluconate

Product identification

Medicine name:
Calcitat N 25 pro inf.

Active substance:
Calcium gluconate monohydrate
Magnesium chloride hexahydrate
Phosphorylcolamine
Calcium hydroxide
Calcium borogluconate

Target species:
Cattle
Cattle (calf)
Dog
Goat
Sheep
Cat
Pig
Pig (piglet)

Route of administration:
Intravenous use

Product details

Active substance and strength:
Calcium gluconate monohydrate
1.55 gram(s) / 100.00 millilitre(s)

Magnesium chloride hexahydrate
3.25 gram(s) / 100.00 millilitre(s)

Phosphorylcolamine
0.30 gram(s) / 100.00 millilitre(s)

Calcium hydroxide
0.66 gram(s) / 100.00 millilitre(s)

Calcium borogluconate
21.45 gram(s) / 100.00 millilitre(s)

Pharmaceutical form:
Solution for infusion

Withdrawal period by route of administration:
Intravenous use:

- **Cattle**
  - Meat and offal. 0 day
- **Cattle (calf)**
  - Meat and offal. 0 day
- **Goat**
  - Meat and offal. 0 day
- **Sheep**

- Meat and offal. 0 day

- **Pig**
  - Meat and offal. 0 day

- **Pig (piglet)**
  - Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QA12AA20

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Germany

Available in:
Germany

Package description:
Available only in German
Available only in German
Available only in German
Available only in German
Available only in German
Available only in German

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:
aniMedica GmbH

Marketing authorisation date:
30/03/1981

Manufacturing sites for batch release:
Pasteur Filiala Filipesti S.A.

Responsible authority:
Federal Office Of Consumer Protection And Food Safety

Authorisation number:
1468.00.00

Date of authorisation status change:
9/11/2009

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Combined File of all Documents

This document does not exist in this language (English). You can find it in another language below.