Calcitat - Injektionslösung für Tiere

Authorised

- Calcium gluconate monohydrate
- Calcium oxide
- Magnesium chloride hexahydrate
- Phosphorylcolamine

Product identification

Medicine name:
Calcitat - Injektionslösung für Tiere

Active substance:
Calcium gluconate monohydrate
Calcium oxide
Magnesium chloride hexahydrate
Phosphorylcolamine

Target species:
Cattle
Dog
Goat
Sheep
Cat
Pig

Route of administration:

Intramuscular use
Intravenous use
Subcutaneous use

Product details

Active substance and strength:
Calcium gluconate monohydrate
215.00 milligram(s) / 1.00 millilitre(s)

Calcium oxide
5.00 milligram(s) / 1.00 millilitre(s)

Magnesium chloride hexahydrate
32.50 milligram(s) / 1.00 millilitre(s)

Phosphorylcolamine
3.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:
Intramuscular use:

- **Cattle**
  - Meat and offal. 0 day
  - Milk. 0 hour
- **Goat**
  - Meat and offal. 0 day
  - Milk. 0 hour
- **Sheep**
  - Meat and offal. 0 day
  - Milk. 0 hour

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

Intravenous use:

- **Cattle**
  - Meat and offal. 0 day
  - Milk. 0 hour
- **Goat**
  - Meat and offal. 0 day
  - Milk. 0 hour
- **Sheep**
  - Meat and offal. 0 day
  - Milk. 0 hour
- **Pig**
  - Meat and offal. 0 day

Subcutaneous use:

- **Cattle**
  - Meat and offal. 0 day
  - Milk. 0 hour
- **Goat**
  - Meat and offal. 0 day
  - Milk. 0 hour
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Sheep

- Meat and offal. 0 day
- Milk. 0 hour

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Pig

- 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:
Austria

Package description:
Available only in German
Available only in German

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Well-established use application (Article 13a of Directive No 2001/82/EC)

Marketing authorisation holder:
aniMedica GmbH

Marketing authorisation date:
3/06/1987

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

Responsible authority:
Austrian Agency For Health And Food Safety

Authorisation number:
8-00020

Date of authorisation status change:
3/06/1987

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

Documents

Package Leaflet

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Summary of Product Characteristics

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

Labelling

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