File downloaded on 2026-11-13
Source URL: https://medicines.health.europa.eu/veterinary/en/600000085261

Calcitat forte - Injektionslösung für Tiere

Authorised

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

Product identification

Medicine name:
Calcitat forte - 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
430.00 milligram(s) / 1.00 millilitre(s)

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

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

Phosphorylcolamine
6.00 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
-

Sheep

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

•

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:
8-00021

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

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

Summary of Product Characteristics

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

Labelling

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