Calcium S 50 aniMedica, otopina za injekciju, za goveda, ovce, koze, svinje, pse i mačke

Authorised

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

Product identification

Medicine name:
Calcium S 50 aniMedica, otopina za injekciju, za goveda, ovce, koze, svinje, pse i mačke

Active substance:
Phosphorylcolamine

Magnesium chloride hexahydrate

Calcium borogluconate

Calcium hydroxide

Calcium gluconate

Target species:
Cattle
Sheep
Goat

Pig
Dog
Cat

Route of administration:
Intramuscular use
Subcutaneous use

Product details

Active substance and strength:
Phosphorylcolamine
6.00 milligram(s) / 1.00 millilitre(s)

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

Calcium borogluconate
429.00 milligram(s) / 1.00 millilitre(s)

Calcium hydroxide
13.20 milligram(s) / 1.00 millilitre(s)

Calcium gluconate
31.00 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
- **Sheep**
  - Meat and offal. 0 day
-

Goat

- Meat and offal. 0 day

•

Pig

- Meat and offal. 0 day

Subcutaneous use:

•

Cattle

- Meat and offal. 0 day

•

Sheep

- Meat and offal. 0 day

•

Goat

- Meat and offal. 0 day

•

Pig

- Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QA12AX

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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Authorisation status:

Valid

Authorised in:

Croatia

Available in:

Croatia

Package description:
Available only in Croatian
Available only in Croatian
Available only in Croatian
Available only in Croatian
Available only in Croatian
Available only in Croatian
Available only in Croatian

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
aniMedica GmbH

Marketing authorisation date:
30/10/2019

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

Responsible authority:
Ministry Of Agriculture Veterinary And Food Safety Directorate

Authorisation number:
UP/I-322-05/13-01/453

Date of authorisation status change:
11/04/2024

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

Documents

Summary of Product Characteristics

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