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Calcitat 25, solution for infusion and injection in cattle

Authorised

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

Product identification

Medicine name:
Calcitat 25, solution for infusion and injection in cattle

Active substance:
Calcium gluconate monohydrate

Calcium borogluconate

Calcium hydroxide

Magnesium chloride hexahydrate

Target species:
Cattle

Route of administration:
Intravenous use

Subcutaneous use

Product details

Active substance and strength:

Calcium gluconate monohydrate
1.55 gram(s) / 100.00 millilitre(s)

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

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

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

Pharmaceutical form:

Solution for injection/infusion

Withdrawal period by route of administration:

Intravenous use:

- **Cattle**
  - Meat and offal. 0 day
  - Milk. 0 day

Subcutaneous use:

- **Cattle**
  - Meat and offal. 0 day
  - Milk. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QA12AA

Legal status of supply:

Veterinary medicinal product not subject to veterinary prescription

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Authorisation status:

Valid

Authorised in:
Ireland

Available in:
Ireland

Package description:
20 500 ml bottlesVial: Sterile plastic bottle (Type PP28).Stopper: Chlorobutyl rubber stopper.Cap: Aluminium cap.
250 ml.Vial: Sterile plastic bottle (Type PP28).Stopper: Chlorobutyl rubber stopper.Cap: Aluminium cap.
1 500 ml bottle.Vial: Sterile plastic bottle (Type PP28).Stopper: Chlorobutyl rubber stopper.Cap: Aluminium cap.

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:
1/10/1989

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

Responsible authority:
Health Products Regulatory Authority

Authorisation number:
VPA10826/001/001

Date of authorisation status change:
1/10/1989

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

Documents

Summary of Product Characteristics