

Calcitat 50, solution for infusion and injection in cattle

Authorised

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

Product identification

Medicine name:

Calcitat 50, 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
3.10 gram(s) / 100.00 millilitre(s)

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

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

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

Pharmaceutical form:

Solution for injection/infusion

Withdrawal period by route of administration:

Intravenous use:

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Cattle

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

Subcutaneous use:

-

Cattle

- Meat and offal. 0 day
 - Milk. 0 day
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Anatomical therapeutic chemical veterinary (ATCvet) codes:

QA12AA

Legal status of supply:

Veterinary medicinal product not subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Ireland

Package description:

12 100 ml bottles.Vial: 100 ml HK2 colourless glass bottle.Stopper: Bromobutyl rubber stopper.Cap: Aluminium cap.

60 250 ml bottles.Vial: 250 ml sterile plastic bottle (Type PP28).Stopper: Chlorobutyl rubber stopper.Cap: Aluminium cap.

1 100 ml bottleVial: 100 ml HK2 colourless glass bottle.Stopper: Bromobutyl rubber stopper.Cap: Aluminium cap.

1 250 ml bottle.Vial: 250 ml sterile plastic bottle (Type PP28).Stopper: Chlorobutyl rubber stopper.Cap: Aluminium cap.

1 500 ml bottle.Vial: 500 ml sterile plastic bottle (Type PP28).Stopper: Chlorobutyl rubber stopper.Cap: Aluminium cap.

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Complete application (stand-alone) - Council Directive 81/851/EEC

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:

VPA10826/001/002

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