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TAT CALCI 50, Solução Injetável para bovinos, equinos, suínos, ovinos e caprinos e cães

Authorised

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

Product identification

Medicine name:
TAT CALCI 50, Solução Injetável para bovinos, equinos, suínos, ovinos e caprinos e cães

Active substance:
Magnesium chloride
Calcium hydroxide
Calcium borogluconate
Calcium gluconate monohydrate

Target species:
Dog
Cattle
Horse
Pig
Sheep

Goat

Route of administration:
Intramuscular use
Intravenous use
Subcutaneous use

Product details

Active substance and strength:
Magnesium chloride
6.50 gram(s) / 100.00 millilitre(s)

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

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

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

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:
Intravenous use:

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

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

Subcutaneous use:

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QA12AA20

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

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Authorisation status:

Valid

Authorised in:
Portugal

Available in:
Portugal

Package description:
Available only in Portuguese
Available only in Portuguese

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:
3/06/1987

Manufacturing sites for batch release:
aniMedica GmbH

Responsible authority:
Directorate General For Food And Veterinary

Authorisation number:
740/01/13NFVPT

Date of authorisation status change:
1/09/2021

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

Documents

Combined File of all Documents

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