MAGNESIO CALCIQUE SOLUTION INJECTABLE

Authorised

- Calcium borogluconate
- Glucose
- Magnesium chloride hexahydrate

Product identification

Medicine name:
MAGNESIO CALCIQUE SOLUTION INJECTABLE

Active substance:
Calcium borogluconate

Glucose

Magnesium chloride hexahydrate

Target species:
Cattle
Pig
Sheep
Goat
Dog

Route of administration:
Intramuscular use
Subcutaneous use
Intravenous use

Intraperitoneal use

Product details

Active substance and strength:

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

Glucose
48.00 milligram(s) / 1.00 millilitre(s)

Magnesium chloride hexahydrate
60.00 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 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

Intravenous 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

Intraperitoneal 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:
QA12AX

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

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Authorisation status:
Valid

Authorised in:
France

Package description:
Available only in French

Available only in French
Available only in French

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:
Dopharma France S.A.S.

Marketing authorisation date:
12/11/1985

Manufacturing sites for batch release:
Dopharma France

Responsible authority:
French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:
FR/V/0912228 0/1985

Date of authorisation status change:
12/11/2010

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

Documents

Summary of Product Characteristics

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Package Leaflet and Labelling

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