CALFOSET soluţie injectablilă

Authorised

- CALCIUM GLYCEROPHOSPHATE
- Calcium gluconate
- Magnesium chloride hexahydrate

Product identification

Medicine name:
CALFOSET soluţie injectablilă

Active substance:
CALCIUM GLYCEROPHOSPHATE
Calcium gluconate
Magnesium chloride hexahydrate

Target species:
Horse
Cattle
Sheep
Goat
Pig

Route of administration:
Intravenous use
Intramuscular use
Subcutaneous use

Product details

Active substance and strength:
CALCIUM GLYCEROPHOSPHATE
8.13 gram(s) / 100.00 millilitre(s)

Calcium gluconate
32.80 gram(s) / 100.00 millilitre(s)

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

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:
Intravenous use:

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

- **Pig**
  - Meat and offal. 0 day

Intramuscular use:

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

Subcutaneous use:

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

Goat

- Meat and offal. 0 day
- Milk. 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

Authorisation status:
Valid

Authorised in:
Romania

Available in:
Romania

Package description:
Available only in Romanian
Available only in Romanian

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:
KRKA tovarna zdravil d.d. Novo mesto

Marketing authorisation date:
18/07/2005

Manufacturing sites for batch release:
KRKA tovarna zdravil d.d. Novo mesto

Responsible authority:
Institute For Control Of Biological Products And Veterinary Medicines

Authorisation number:
110325

Date of authorisation status change:
17/11/2011

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.