CALFOSET injekčný roztok pre kone, hovädzí dobytok, ovce, kozy, ošípané a odstavčatá

Authorised

- Calcium gluconate monohydrate
- CALCIUM GLYCEROPHOSPHATE
- Magnesium chloride hexahydrate

Product identification

Medicine name:
CALFOSET injekčný roztok pre kone, hovädzí dobytok, ovce, kozy, ošípané a odstavčatá

Active substance:
Calcium gluconate monohydrate
CALCIUM GLYCEROPHOSPHATE
Magnesium chloride hexahydrate

Target species:
Cattle
Sheep
Goat
Pig
Pig (weaned piglet)
Horse

Route of administration:

Subcutaneous use
Intravenous use
Intramuscular use

Product details

Active substance and strength:
Calcium gluconate monohydrate
328.20 milligram(s) / 1.00 millilitre(s)

CALCIUM GLYCEROPHOSPHATE
81.30 milligram(s) / 1.00 millilitre(s)

Magnesium chloride hexahydrate
41.80 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:
Subcutaneous use:

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

- Meat and offal. 0 day

Intravenous use:

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

Intramuscular use:

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

- Milk. 0 hour

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

- **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:
Slovakia

Available in:
Slovakia

Package description:
Available only in Slovak
Available only in Slovak

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:
28/03/1995

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

Responsible authority:
Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:
96/0014/95-S

Date of authorisation status change:
28/03/1995

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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