Surcalce - Injektionslösung für Pferd, Rind, Schwein, Schaf, Ziege

Authorised

- Calcium gluconate monohydrate
- Calcium acetate
- Magnesium hypophosphite hexahydrate

Product identification

Medicine name:
Surcalce - Injektionslösung für Pferd, Rind, Schwein, Schaf, Ziege

Active substance:
Calcium gluconate monohydrate
Calcium acetate
Magnesium hypophosphite hexahydrate

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

Route of administration:
Intravenous use
Subcutaneous use

Product details

Active substance and strength:

Calcium gluconate monohydrate
465.00 milligram(s) / 1.00 millilitre(s)

Calcium acetate
37.00 milligram(s) / 1.00 millilitre(s)

Magnesium hypophosphite hexahydrate
30.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Intravenous use:

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

Subcutaneous use:

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QA12AA20

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

Authorisation status:
Valid

Authorised in:
Austria

Package description:
Available only in German

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:
V.M.D.

Marketing authorisation date:
4/12/1978

Manufacturing sites for batch release:
Sanochemia Pharmazeutika AG

Responsible authority:
Austrian Agency For Health And Food Safety

Authorisation number:
16368

Date of authorisation status change:
4/12/1978

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

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Labelling

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