SURCALCE injekčný roztok

Authorised

- Calcium gluconate
- Calcium acetate
- Magnesium hypophosphite hexahydrate

Product identification

Medicine name:
SURCALCE injekčný roztok

Active substance:
Calcium gluconate
Calcium acetate
Magnesium hypophosphite hexahydrate

Target species:
Cattle
Horse
Sheep
Goat
Pig

Route of administration:
Subcutaneous use
Intravenous use

Product details

Active substance and strength:

Calcium gluconate
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:

Subcutaneous use:

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

- Milk. 0 day Zero days
- Meat and offal. 0 day Zero days

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Pig

- Meat and offal. 0 day Zero days

Intravenous use:

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Cattle

- Milk. 0 day Zero days
- Meat and offal. 0 day Zero days

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Horse

- Milk. 0 day Zero days
- Meat and offal. 0 day Zero days

•

Sheep

- Milk. 0 day Zero days
- Meat and offal. 0 day Zero days

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Goat

- Milk. 0 day Zero days
- Meat and offal. 0 day Zero days

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Pig

- Meat and offal. 0 day Zero days

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QA12AX

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

Authorisation status:
Valid

Authorised in:
Slovakia

Package description:
Available only in Slovak

Additional information

Entitlement type:
Marketing Authorisation

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

Marketing authorisation holder:
V.M.D.

Marketing authorisation date:
23/12/1994

Manufacturing sites for batch release:
Sanochemia Pharmazeutika AG

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

Authorisation number:

96/398/91-S

Date of authorisation status change:
23/12/1994

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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