

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:**

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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

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

Labelling

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