Ca-mg-infuus, oplossing voor infusie voor runderen en schapen.

Authorised

- Calcium gluconate
- Magnesium chloride
- Calcium oxide

Product identification

Medicine name:
Ca-mg-infuus, oplossing voor infusie voor runderen en schapen.

Active substance:
Calcium gluconate

Magnesium chloride

Calcium oxide

Target species:
Cattle
Sheep

Route of administration:
Intravenous use
Subcutaneous use

Product details

Active substance and strength:

Calcium gluconate
119.30 milligram(s) / 1.00 millilitre(s)

Magnesium chloride
37.10 milligram(s) / 1.00 millilitre(s)

Calcium oxide
7.60 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for infusion

Withdrawal period by route of administration:

Intravenous use:

- **Cattle**
  - Meat and offal. no withdrawal period — Withdrawal period is zero days
  - Milk. no withdrawal period — Withdrawal period is zero days
  - Milk. no withdrawal period — Withdrawal period is zero days
- **Sheep**
  - Meat and offal. no withdrawal period — Withdrawal period is zero days

Subcutaneous use:

- **Cattle**
  - Meat and offal. no withdrawal period — Withdrawal period is zero days
  - Milk. no withdrawal period — Withdrawal period is zero days
- **Sheep**
  - Meat and offal. no withdrawal period — Withdrawal period is zero days

- Milk. no withdrawal period	Withdrawal period is 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:
Netherlands

Available in:
Netherlands

Package description:
Available only in Dutch
Available only in Dutch

Additional information

Entitlement type:
Marketing Authorisation

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

Marketing authorisation holder:
Eurovet Animal Health B.V.

Marketing authorisation date:
16/01/1992

Manufacturing sites for batch release:
Eurovet Animal Health B.V.

Responsible authority:
Medicines Evaluation Board

Authorisation number:
REG NL 1609

Date of authorisation status change:
28/05/2014

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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