

# 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](http://www.adrreports.eu/vet)

## Documents

Combined File of all Documents

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