# Calmasol-440, solution for infusion for cattle, sheep and pigs

Authorised

- Calcium gluconate monohydrate
- Magnesium chloride hexahydrate
- Boric acid

## Product identification

**Medicine name:**
Calmasol-440, solution for infusion for cattle, sheep and pigs
Calmafusion vet 380 mg/ml + 60 mg/ml + 50 mg/ml infusjonsvæske, oppløsning til storfe, sau og gris

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**Active substance:**
Calcium gluconate monohydrate

Magnesium chloride hexahydrate

Boric acid

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**Target species:**
Cattle
Sheep
Pig

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**Route of administration:**
Intravenous use

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## Product details

**Active substance and strength:**

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

Magnesium chloride hexahydrate
60.00 milligram(s) / 1.00 millilitre(s)

Boric acid
50.00 milligram(s) / 1.00 millilitre(s)

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**Pharmaceutical form:**

Solution for infusion

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**Withdrawal period by route of administration:**

**Intravenous use:**

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

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QA12AX

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Norway

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**Package description:**
Graduated polypropylene bottle, closed with a bromobutyl rubber stopper and secured with an aluminium cap. 12x500 ml.
Graduated polypropylene bottle, closed with a bromobutyl rubber stopper and secured with an aluminium cap. 1x500 ml.

## Additional information

**Entitlement type:**
Marketing Authorisation

**Legal basis of product authorisation:**
Generic application (Article 13(1) of Directive No 2001/82/EC)

**Marketing authorisation holder:**
Interchemie Werken De Adelaar Eesti AS

**Marketing authorisation date:**
17/12/2019

**Manufacturing sites for batch release:**
Interchemie Werken De Adelaar Eesti AS

**Responsible authority:**
Norwegian Medicines Agency

**Authorisation number:**
19-12901

**Date of authorisation status change:**
17/12/2019

**Reference member state:**
Estonia

**Procedure number:**
EE/V/0104/001

**Concerned member states:**
Austria Belgium Bulgaria Croatia Cyprus Czechia Denmark Finland France Germany Greece Hungary Iceland Ireland Italy Latvia Malta Netherlands Norway Poland Portugal Romania Slovakia Slovenia Spain Sweden United Kingdom (Northern Ireland)

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

## Documents

Summary of Product Characteristics

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