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# Calcibel Forte, 380/60/50 mg/ml solution for infusion for horses, cattle, sheep, goats and pigs

Authorised

- Calcium gluconate
- Magnesium chloride hexahydrate
- Boric acid

## Product identification

**Medicine name:**
Calcibel Forte, 380/60/50 mg/ml solution for infusion for horses, cattle, sheep, goats and pigs

**Active substance:**
Calcium gluconate
Magnesium chloride hexahydrate
Boric acid

**Target species:**
Horse
Cattle
Sheep
Goat
Pig

**Route of administration:**
Intravenous use

## Product details

**Active substance and strength:**

Calcium gluconate
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:**

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

- **Pig**
  - Meat and offal. 0 day

**Anatomical therapeutic chemical veterinary (ATCvet) codes:**
QA12AX

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

**Authorisation status:**
Valid

**Authorised in:**
Hungary

**Available in:**
Hungary

**Package description:**
Plastic Bottle 12 x 500.0 millilitre(s)
Plastic Bottle 1 x 500.0 millilitre(s)

## 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:**
Bela-Pharm GmbH & Co. KG

**Marketing authorisation date:**
19/10/2021

**Manufacturing sites for batch release:**

Bela-Pharm GmbH & Co. KG

**Responsible authority:**
Directorate Of Veterinary Medicinal Products

**Authorisation number:**
4285/X/21 NÉBIH ÁTI

**Date of authorisation status change:**
19/10/2021

**Reference member state:**
Czechia

**Procedure number:**
CZ/V/0170/001

**Concerned member states:**
Austria Cyprus Denmark Estonia Finland Greece Hungary Iceland Ireland Italy Latvia Lithuania Poland Portugal Romania Slovakia Slovenia Spain United Kingdom (Northern Ireland)

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

## Documents

Summary of Product Characteristics

Package Leaflet

Labelling