

# 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

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### **Active substance:**

Calcium gluconate

Magnesium chloride hexahydrate

Boric acid

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### **Target species:**

Horse

Cattle

Sheep

Goat

Pig

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### **Route of administration:**

Intravenous use

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## 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:**

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#### **Horse**

- Milk. 0 hour

- Meat and offal. 0 day

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#### **Cattle**

- Milk. 0 hour

- Meat and offal. 0 day

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#### **Sheep**

- Milk. 0 hour

- Meat and offal. 0 day

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#### **Goat**

- Milk. 0 hour

- Meat and offal. 0 day

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**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:**

Latvia

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**Available in:**

Latvia

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**Package description:**

Plastic Bottle 12 x 500.0 millilitre(s)

Plastic Bottle 1 x 500.0 millilitre(s)

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Generic application (Article 13(1) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Bela-Pharm GmbH & Co. KG

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**Marketing authorisation date:**

23/09/2021

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**Manufacturing sites for batch release:**

Bela-Pharm GmbH & Co. KG

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**Responsible authority:**

Food And Veterinary Service

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**Authorisation number:**

V/DCP/21/0049

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**Date of authorisation status change:**

23/09/2021

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**Reference member state:**

Czechia

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**Procedure number:**

CZ/V/0170/001

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**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)

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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.

Summary of Product Characteristics

Package Leaflet

Labelling