

# 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

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

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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:**

Ireland

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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:**

12/11/2021

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**Manufacturing sites for batch release:**

Bela-Pharm GmbH & Co. KG

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**Responsible authority:**

Health Products Regulatory Authority

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**Authorisation number:**

VPA10445/007/001

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**Date of authorisation status change:**

12/11/2021

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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)