

# CALFORMAG, Injekční roztok

Not authorised

- PHOSPHORUS
- CALCIUM
- MAGNESIUM

## Product identification

**Medicine name:**

CALFORMAG, Injekční roztok

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**Active substance:**

PHOSPHORUS

CALCIUM

MAGNESIUM

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**Target species:**

Cattle

Horse

Sheep

Goat

Pig

Dog

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**Route of administration:**

Intravenous use

Subcutaneous use

Intramuscular use

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## Product details

### **Active substance and strength:**

PHOSPHORUS

10.10 milligram(s) / 1.00 millilitre(s)

CALCIUM

44.60 milligram(s) / 1.00 millilitre(s)

MAGNESIUM

6.00 milligram(s) / 1.00 millilitre(s)

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### **Pharmaceutical form:**

Solution for injection

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### **Withdrawal period by route of administration:**

#### **Intravenous use:**

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#### **Cattle**

- Meat. 0 day

- Milk. 0 day

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#### **Horse**

- Meat. 0 day

- 

#### **Sheep**

- Meat. 0 day

- Milk. 0 day

- 

#### **Goat**

- Meat. 0 day

- Milk. 0 day

-

**Pig**

- Meat. 0 day

**Subcutaneous use:**

•

**Cattle**

- Meat. 0 day

- Milk. 0 day

•

**Horse**

- Meat. 0 day

•

**Sheep**

- Meat. 0 day

- Milk. 0 day

•

**Goat**

- Meat. 0 day

- Milk. 0 day

•

**Pig**

- Meat. 0 day

**Intramuscular use:**

•

**Cattle**

- Meat. 0 day

- Milk. 0 day

•

**Horse**

- Meat. 0 day

- 

**Sheep**

- Meat. 0 day
- Milk. 0 day

- 

**Goat**

- Meat. 0 day
- Milk. 0 day

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**Pig**

- Meat. 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:**

Surrendered

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**Authorised in:**

Czechia

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**Package description:**

Available only in Czech

Available only in Czech

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Full application (Article 12(3) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

BB Pharma a.s.

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**Marketing authorisation date:**

2/01/2002

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**Manufacturing sites for batch release:**

Farmacia Martin a.s.

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**Responsible authority:**

Institute For State Control Of Veterinary Biologicals And Medicines

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**Authorisation number:**

96/002/02-C

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**Date of authorisation status change:**

19/10/2022

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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

Summary of Product Characteristics

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

Package Leaflet

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

Labelling

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