

# Calcibel 240/60/60 mg/ml solution for infusion for horses, cattle, sheep, goats and pigs

Authorised

- Boric acid
- Calcium gluconate
- Magnesium chloride hexahydrate

## Product identification

### **Medicine name:**

Calcibel 240/60/60 mg/ml solution for infusion for horses, cattle, sheep, goats and pigs

### **Active substance:**

Boric acid

Calcium gluconate

Magnesium chloride hexahydrate

### **Target species:**

Cattle

Sheep

Horse

Pig

Goat

### **Route of administration:**

Intravenous use

## Product details

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

Boric acid

60.00 milligram(s) / 1.00 millilitre(s)

Calcium gluconate

240.00 milligram(s) / 1.00 millilitre(s)

Magnesium chloride hexahydrate

60.00 milligram(s) / 1.00 millilitre(s)

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### **Pharmaceutical form:**

Solution for infusion

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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:**

Spain

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### **Available in:**

Spain

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### **Package description:**

Bottle for infusion made of polypropylene with scaling, closures made of bromobutyl rubber stopper, aluminium caps with tear-off. Pack size 1 x 500 ml.

Bottle for infusion made of polypropylene with scaling, closures made of bromobutyl rubber stopper, aluminium caps with tear-off. Pack size 12 x 500 ml.

Bottle for infusion made of polypropylene with scaling, closures made of bromobutyl rubber stopper, aluminium caps with tear-off. Pack size 6 x 500 ml.

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Hybrid application (Article 13(3) 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:**

6/05/2016

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**Manufacturing sites for batch release:**

Bela-Pharm GmbH & Co. KG

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**Responsible authority:**

Spanish Agency Of Medicines And Medical Devices

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**Authorisation number:**

3398 ESP

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**Date of authorisation status change:**

18/02/2021

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**Reference member state:**

Netherlands

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**Procedure number:**

NL/V/0197/001

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**Concerned member states:**

Austria Croatia Cyprus Czechia Denmark Finland Greece Hungary Ireland  
Poland Portugal Romania Slovakia Slovenia Spain Sweden  
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

### 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.