# 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
CALZIUM 240/60/60 mg/ml SOLUCION PARA PERFUSION PARA CABALLOS, BOVINO, OVINO,CAPRINO Y PORCINO

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

**Target species:**
Cattle
Goat (adult female)
Sheep
Horse
Pig

**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)

**Pharmaceutical form:**
Solution for infusion

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

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

**Authorisation status:**
Valid

**Authorised in:**
Spain

**Available in:**
Spain

**Package description:**
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.
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 1 x 500 ml.

## Additional information

**Entitlement type:**
Marketing Authorisation

**Legal basis of product authorisation:**
Hybrid application (Article 13(3) of Directive No 2001/82/EC)

**Marketing authorisation holder:**
Bela-Pharm GmbH & Co. KG

**Marketing authorisation date:**
6/05/2016

**Manufacturing sites for batch release:**
Bela-Pharm GmbH & Co. KG

**Responsible authority:**
Spanish Agency For Medicines And Health Products

**Authorisation number:**
3398 ESP

**Date of authorisation status change:**
18/02/2021

**Reference member state:**
Netherlands

**Procedure number:**
NL/V/0197/001

**Concerned member states:**
Austria Croatia Cyprus Czechia Denmark Finland Greece Hungary Ireland Poland Portugal Romania Slovakia Slovenia Spain Sweden

United Kingdom (Northern Ireland)

To consult adverse reactions on veterinary medicinal products please go to 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.