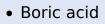
# Calcibel 240/60/60 mg/ml solution for infusion for horses, cattle, sheep, goats and pigs

Authorised



- 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

Calcibel 240/60/60 mg/ml infúzny roztok pre kone, hovädzí dobytok, ovce, kozy a ošípané

#### **Active substance:**

Boric acid Calcium gluconate Magnesium chloride hexahydrate

## **Target species:**

Cattle Goat (adult female) Sheep Horse Pig

**Route of administration:** 

# **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

#### Withdrawal period by route of administration:

#### Intravenous use:

```
•
Cattle
•
Goat (adult female)
•
Sheep
•
Horse
•
Pig
```

## Anatomical therapeutic chemical veterinary (ATCvet) codes:

QA12AX

## Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

#### Authorisation status:

Valid

## Authorised in:

Slovakia

## Available in:

Slovakia

#### 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:

1/07/2016

## Manufacturing sites for batch release:

Bela-Pharm GmbH & Co. KG

## Responsible authority:

Institute For State Control Of Veterinary Biologicals And Medicaments

# Authorisation number: 96/033/DC/16-S

# Date of authorisation status change:

1/07/2016

#### **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 <u>www.adrreports.eu/vet</u>

# Documents

Combined File of all Documents

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