

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

Calcibel 240/60/60 mg/ml raztopina za infundiranje za konje, govedo, ovce, koze in prašiče

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:

Slovenia

Available in:

Slovenia

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:

22/03/2016

Manufacturing sites for batch release:

Bela-Pharm GmbH & Co. KG

Responsible authority:

Agency For Medicinal Products And Medical Devices Of The Republic Of Slovenia

Authorisation number:

DC/V/0527/001

Date of authorisation status change:

22/03/2016

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 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.

Package Leaflet and Labelling

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