

Calciolab 216.18/60/51 mg/ml solution for infusion

Authorised

- Calcium gluconate
- Magnesium chloride hexahydrate
- Magnesium hypophosphite hexahydrate

Product identification

Medicine name:

Calciolab 216.18/60/51 mg/ml solution for infusion

Calciolab 216,18/60/51 mg/ml roztok na infúziu

Active substance:

Calcium gluconate

Magnesium chloride hexahydrate

Magnesium hypophosphite hexahydrate

Target species:

Cattle

Sheep

Goat

Pig

Horse

Dog

Route of administration:

Intravenous use

Product details

Active substance and strength:

Calcium gluconate

216.18 milligram(s) / 1.00 millilitre(s)

Magnesium chloride hexahydrate

60.00 milligram(s) / 1.00 millilitre(s)

Magnesium hypophosphite hexahydrate

51.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for infusion

Withdrawal period by route of administration:

Intravenous use:

-

Cattle

- Meat and offal. 0 day
- Milk. 0 day

-

Sheep

- Meat and offal. 0 day
- Milk. 0 day

-

Goat

- Meat and offal. 0 day
- Milk. 0 day

-

Pig

- Meat and offal. 0 day

-

Horse

- Meat and offal. 0 day
- Milk. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QA12

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Slovakia

Package description:

Polypropylene bottles with chlorobutyl rubber stopper and aluminium cap. Box with 1 bottle of 100 ml.

Polypropylene bottles with chlorobutyl rubber stopper and aluminium cap. Bottle of 500 ml.

Polypropylene bottles with chlorobutyl rubber stopper and aluminium cap. Box with 10 bottles of 100 ml.

Polypropylene bottles with chlorobutyl rubber stopper and aluminium cap. Box with 6 bottles of 500 ml.

Polypropylene bottles with chlorobutyl rubber stopper and aluminium cap. Box with 12 bottles of 500 ml.

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Generic application (Article 18 of Regulation (EU) 2019/6)

Marketing authorisation holder:

Labiana Life Sciences S.A.

Marketing authorisation date:

4/12/2024

Manufacturing sites for batch release:

Labiana Life Sciences S.A.

Responsible authority:

Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:

96/056/DC/24-S

Date of authorisation status change:

4/12/2024

Reference member state:

Estonia

Procedure number:

EE/V/0107/001

Concerned member states:

Bulgaria Czechia Greece Hungary Latvia Lithuania Slovakia

Generic of:

600000055967

To consult adverse reactions on veterinary medicinal products please go to
www.adrreports.eu/vet

Documents

Combined File of all Documents

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