

Calmagluc (60 mg + 22 mg + 30 mg + 100 mg)/ml Roztwór do wstrzykiwań

Authorised

- Glucose
- Calcium gluconate
- Calcium hypophosphite
- Magnesium chloride hexahydrate

Product identification

Medicine name:

Calmagluc (60 mg + 22 mg + 30 mg + 100 mg)/ml Roztwór do wstrzykiwań

Active substance:

Glucose

Calcium gluconate

Calcium hypophosphite

Magnesium chloride hexahydrate

Target species:

Cattle

Horse

Pig

Dog

Route of administration:

Intramuscular use

Intravenous use

Product details

Active substance and strength:

Glucose

100.00 milligram(s)/millilitre / 1.00 milligram(s)/millilitre

Calcium gluconate

60.00 milligram(s)/millilitre / 1.00 milligram(s)/millilitre

Calcium hypophosphite

22.00 milligram(s)/millilitre / 1.00 milligram(s)/millilitre

Magnesium chloride hexahydrate

30.00 milligram(s)/millilitre / 1.00 milligram(s)/millilitre

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Intramuscular use:

-

Cattle

- Milk. 0 hour

- Meat and offal. 0 day

-

Horse

- Meat and offal. 0 day

-

Pig

- Meat and offal. 0 day

Intravenous use:

-

Cattle

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

-

Horse

- Meat and offal. 0 day

-

Pig

- Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QA12AX

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Poland

Package description:

Available only in Polish

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Well-established use application (Article 13a of Directive No 2001/82/EC)

Marketing authorisation holder:

Biowet Pulawy Sp. z o.o.

Marketing authorisation date:

18/09/2002

Manufacturing sites for batch release:

Biowet Pulawy Sp. z o.o.

Responsible authority:

Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

Authorisation number:

1317

Date of authorisation status change:

18/09/2002

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

Documents

Package Leaflet

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Summary of Product Characteristics

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.