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Kaltetan, 250 mg/ml + 80 mg/ml + 10 mg/ml solution for infusion for horses, cattle and pigs

Authorised

- CALCIUM GLUCONATE FOR INJECTION
- Magnesium chloride hexahydrate
- Sodium glycerophosphate pentahydrate

Product identification

Medicine name:
Kaltetan, 250 mg/ml + 80 mg/ml + 10 mg/ml solution for infusion for horses, cattle and pigs

Active substance:
CALCIUM GLUCONATE FOR INJECTION
Magnesium chloride hexahydrate
Sodium glycerophosphate pentahydrate

Target species:
Horse
Pig
Cattle

Route of administration:
Intravenous use

Product details

Active substance and strength:

CALCIUM GLUCONATE FOR INJECTION
250.00 milligram(s)/millilitre / 1.00 milligram(s)/millilitre

Magnesium chloride hexahydrate
80.00 milligram(s) / 1.00 millilitre(s)

Sodium glycerophosphate pentahydrate
10.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for infusion

Withdrawal period by route of administration:

Intravenous use:

- **Horse**
  - Milk. 5 week — Meat and offal: Zero days. Milk: Zero hours
- **Pig**
  - Meat and offal. 5 week — Meat and offal: Zero days
- **Cattle**
  - Milk. 5 week — Meat and offal: Zero days. Milk: Zero hours

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Anatomical therapeutic chemical veterinary (ATCvet) codes:

QA12AX

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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Authorisation status:

Valid

Authorised in:
Poland

Available in:
Poland

Package description:
Polypropylene (PP) bottles closed with a bromobutyl rubber stopper, type I and secured with an aluminium cap. Package size: 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:
Przedsiebiorstwo Wielobranzowe Vet-Agro Sp. z o.o.

Marketing authorisation date:
7/04/2022

Manufacturing sites for batch release:
Przedsiebiorstwo Wielobranzowe Vet-Agro Sp. z o.o.

Responsible authority:
Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

Authorisation number:
3171

Date of authorisation status change:
7/04/2022

Reference member state:
Poland

Procedure number:

PL/V/0110/001

Concerned member states:
Bulgaria Czechia Greece Hungary Italy Lithuania Portugal Romania Spain

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

Documents

Summary of Product Characteristics

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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.