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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:
Greece

Available in:
Greece

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:
3/04/2022

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

Responsible authority:
National Organization For Medicines

Authorisation number:
34226/04-04-2022/K-0248501

Date of authorisation status change:
3/04/2022

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.