File downloaded on 2026-11-28
Source URL: https://medicines.health.europa.eu/veterinary/en/600000092620

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

---

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QA12AX

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

---

Authorisation status:

Valid

Authorised in:
Romania

Available in:
Romania

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:
15/03/2022

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

Responsible authority:
Institute For Control Of Biological Products And Veterinary Medicines

Authorisation number:
220055

Date of authorisation status change:
10/06/2025

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

Combined File of all Documents

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