

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

Kaltetan, 250 mg/ml + 80 mg/ml + 10 mg/ml soluzione per infusione per cavalli, bovini e suini

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:

Italy

Available in:

Italy

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:

Vet-Agro Multi-Trade Company Sp. z o.o.

Marketing authorisation date:

20/07/2022

Manufacturing sites for batch release:

Multi-Trade Company "Vet-Agro" Sp. z o.o.

Responsible authority:

Ministry Of Health

Authorisation number:

105612

Date of authorisation status change:

20/07/2022

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

Combined File of all Documents

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

Source URL: <https://medicines.health.europa.eu/veterinary/600000092612>