

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

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Pig

- Meat and offal. 5 week Meat and offal: Zero days

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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:

Hungary

Available in:

Hungary

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:

Przedsiębiorstwo Wielobranzowe Vet-Agro Sp. z o.o.

Marketing authorisation date:

8/02/2022

Manufacturing sites for batch release:

Przedsiębiorstwo Wielobranzowe Vet-Agro Sp. z o.o.

Responsible authority:

Directorate Of Veterinary Medicinal Products

Authorisation number:

4308/X/22 NÉBIH ÁTI

Date of authorisation status change:

8/02/2022

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