

# Kaltetan forte, 458.4 mg/ml + 125 mg/ml + 20 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 forte, 458.4 mg/ml + 125 mg/ml + 20 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

Cattle

Pig

### Route of administration:

Intravenous use

## Product details

### Active substance and strength:

CALCIUM GLUCONATE FOR INJECTION

458.40 milligram(s) / 1.00 millilitre(s)

Magnesium chloride hexahydrate

125.00 milligram(s) / 1.00 millilitre(s)

Sodium glycerophosphate pentahydrate

20.00 milligram(s) / 1.00 millilitre(s)

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### Pharmaceutical form:

Solution for infusion

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### Withdrawal period by route of administration:

#### Intravenous use:

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#### Horse

- Milk. 0 week  
Meat and offal: Zero days. Milk: Zero hours

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#### Cattle

- Meat and offal. 0 week  
Meat and offal: Zero days. Milk: Zero hours

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#### Pig

- Meat and offal. 0 week  
Meat and offal: Zero days

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### Anatomical therapeutic chemical veterinary (ATCvet) codes:

QA12AX

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### Legal status of supply:

Veterinary medicinal product not subject to veterinary prescription

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### Authorisation status:

Valid

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**Authorised in:**

Lithuania

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**Available in:**

Lithuania

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**Package description:**

Polypropylene (PP) bottles closed with a bromobutyl rubber stopper, type I and secured with an aluminium cap. Package size: 500 ml

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Hybrid application (Article 13(3) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

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

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**Marketing authorisation date:**

25/01/2022

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**Manufacturing sites for batch release:**

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

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**Responsible authority:**

State Food And Veterinary Service

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**Authorisation number:**

LT/2/22/2697/001

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**Date of authorisation status change:**

27/04/2025

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**Reference member state:**

Poland

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**Procedure number:**

PL/V/0110/002

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**Concerned member states:**

Bulgaria Czechia Greece Hungary Italy Lithuania Portugal Romania Spain

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To consult adverse reactions on veterinary medicinal products please go to  
[www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Combined File of all Documents

English (PDF)

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