

# 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 solução para perfusão para equinos, bovinos e suínos.

Kaltetan, 250 mg/ml + 80 mg/ml + 10 mg/ml solution for infusion for horses, cattle and pigs

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### **Active substance:**

CALCIUM GLUCONATE FOR INJECTION

Magnesium chloride hexahydrate

Sodium glycerophosphate pentahydrate

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### **Target species:**

Horse

Pig

Cattle

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### **Route of administration:**

Intravenous use

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## 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)

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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. 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

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

QA12AX

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### **Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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### **Authorisation status:**

Valid

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**Authorised in:**

Portugal

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**Available in:**

Portugal

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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:**

17/02/2022

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**Manufacturing sites for batch release:**

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

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**Responsible authority:**

Directorate General For Food And Veterinary

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**Authorisation number:**

1495/01/22DFVPT

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**Date of authorisation status change:**

30/04/2025

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**Reference member state:**

Poland

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**Procedure number:**

**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](http://www.adrreports.eu/vet)

## Documents

Combined File of all Documents

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