

# Catosal 100 mg/ml + 0.05 mg/ml solution for injection for cattle, horses and dogs

Authorised

- Butafosfan
- Cyanocobalamin

## Product identification

### **Medicine name:**

Catosal 100 mg/ml + 0.05 mg/ml solution for injection for cattle, horses and dogs

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### **Active substance:**

Butafosfan

Cyanocobalamin

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### **Target species:**

Cattle

Dog

Horse

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### **Route of administration:**

Intravenous use

Intramuscular use

Subcutaneous use

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## Product details

### Active substance and strength:

Butafosfan

100.00 milligram(s) / 1.00 millilitre(s)

Cyanocobalamin

0.05 milligram(s) / 1.00 millilitre(s)

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### Pharmaceutical form:

Solution for injection

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### Withdrawal period by route of administration:

#### Intravenous use:

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#### Cattle

- Meat and offal. 0 day

- Milk. 0 hour

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#### Horse

- Meat and offal. 0 day

- Milk. 0 hour

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

QA12CX99

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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:

Czechia

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### Available in:

Czechia

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**Package description:**

(ID3) 50 millilitre(s): Box (board) with 1 Vial (Glass type II) with 50 millilitre(s), closed with Stopfen (chlorobutyl rubber) and Cap`` (aluminium)

(ID2) 250 millilitre(s): Box (board) with 1 Vial (Glass type I) with 250 millilitre(s), closed with Stopfen (chlorobutyl rubber) and Cap`` (aluminium)

(ID1) 100 millilitre(s): Box (board) with 1 Vial (Glass type II) with 100 millilitre(s), closed with Stopfen (chlorobutyl rubber) and Cap`` (aluminium)

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Legal basis reviewed according to Acquis communautaire

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**Marketing authorisation holder:**

Elanco Animal Health GmbH

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**Marketing authorisation date:**

9/09/1994

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**Manufacturing sites for batch release:**

KVP Pharma+Veterinaer Produkte GmbH

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**Responsible authority:**

Institute For State Control Of Veterinary Biologicals And Medicaments

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**Authorisation number:**

96/985/94-C

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**Date of authorisation status change:**

7/04/2011

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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

Summary of Product Characteristics

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

Package Leaflet

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

Labelling

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