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# 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

**Active substance:**
Butafosfan
Cyanocobalamin

**Target species:**
Cattle
Dog
Horse

**Route of administration:**
Intravenous use
Intramuscular use
Subcutaneous use

## 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:**

- **Cattle**
  - Meat and offal. 0 day
  - Milk. 0 day
- **Horse**
  - Meat and offal. 0 day
  - Milk. 0 day

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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:**
Belgium

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**Available in:**

Belgium

**Package description:**
(ID3) 50 millilitre(s): Box (Cardboard) with 1 Vial (Glass type II) with 50 millilitre(s), closed with Stopper (Chlorobutyl Rubber) and Cap (Aluminium)
(ID2) 250 millilitre(s): Box (Cardboard) with 1 Vial (Glass type I) with 250 millilitre(s), closed with Stopper (Chlorobutyl Rubber) and Cap (Aluminium)
(ID1) 100 millilitre(s): Box (Cardboard) with 1 Vial (Glass type II) with 100 millilitre(s), closed with Stopper (Chlorobutyl Rubber) and Cap (Aluminium)

## Additional information

**Entitlement type:**
Marketing Authorisation

**Legal basis of product authorisation:**
Legal basis reviewed according to Acquis communautaire

**Marketing authorisation holder:**
Elanco Animal Health GmbH

**Marketing authorisation date:**
2/06/1965

**Manufacturing sites for batch release:**
KVP Pharma+Veterinaer Produkte GmbH

**Responsible authority:**
Federal Agency For Medicines And Health Products

**Authorisation number:**
This information is not available for this product.

**Date of authorisation status change:**
24/10/2024

**Reference member state:**
Germany

**Procedure number:**

DE/V/0390/001

**Concerned member states:**
Austria Belgium Bulgaria Croatia Cyprus Czechia France Greece Hungary Italy Luxembourg Netherlands Poland Portugal Romania Slovakia Slovenia Spain

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

## Documents

Labelling

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.

Summary of Product Characteristics

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

Combined File of all Documents