# 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 Solution injectable
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 hour
- **Dog**
- **Horse**
  - Meat and offal. 0 day
  - Milk. 0 hour

**Intramuscular use:**

- **Dog**

**Subcutaneous use:**

- **Dog**

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

QA12CX91

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**Legal status of supply:**
Veterinary medicinal product subject to veterinary prescription

**Authorisation status:**
Valid

**Authorised in:**
Luxembourg

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

## 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:**
1/06/1965

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

**Responsible authority:**
Ministry Of Health And Social Security

**Authorisation number:**
V/442/07/10/0924

**Date of authorisation status change:**
1/06/1965

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

## Documents

Summary of Product Characteristics

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.

Package Leaflet

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

Combined File of all Documents

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