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 Oplossing voor injectie
Catosal 100 mg/ml - 0.05 mg/ml Solution injectable
Catosal 100 mg/ml - 0.05 mg/ml Injektionslösung
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)

Pharmaceutical form:
Solution for injection

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**

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QA12CX99

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

Authorisation status:
Valid

Authorised in:
Belgium

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 Cap and Cap and Stopper (chlorobutyl rubber`, Aluminium, Aluminium, chlorobutyl rubber`)
(ID3) 50 millilitre(s): Box (Cardboard) with 1 Vial (Glass type II) with 50 millilitre(s), closed with Stopper and Cap (chlorobutyl rubber`, 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