Source URL: https://medicines.health.europa.eu/veterinary/en/60000005080

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



- Butafosfan
- Cyanocobalamin

Product identification

Medicine name:

Catosal 100 mg/ml + 0,05 mg/ml raztopina za injiciranje za govedo, konje in pse 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:

Slovenia

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:

1/03/2002

Manufacturing sites for batch release:

KVP Pharma+Veterinaer Produkte GmbH

Responsible authority:

Agency For Medicinal Products And Medical Devices Of The Republic Of Slovenia

| Authorisation number: MR/V/0059/001 |
|--|
| Date of authorisation status change: 17/10/2024 |
| 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. |
| 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. |

| Combined File of all Documents | | |
|--------------------------------|--|--|
| | | |
| | | |