Source URL: https://medicines.health.europa.eu/veterinary/en/600000060182

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



- Butafosfan
- Cyanocobalamin

Product identification

Medicine name:

CATOSAL, 100/0,05mg/ml, Injekční roztok
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

•

Horse

- Meat and offal. 0 day
- Milk. 0 hour

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QA12CX99

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Czechia

Package description:

(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) (ID3) 50 millilitre(s): Box (board) with 1 Vial (Glass type II) with 50 millilitre(s), closed with Stopfen (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:

9/09/1994

Manufacturing sites for batch release:

KVP Pharma+Veterinaer Produkte GmbH

Responsible authority:

Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:

96/985/94-C

Date of authorisation status change:

7/04/2011

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

Documents