

Vey Tosal 100 mg/ml + 0.05 mg/ml solution for injection for horses, cattle, dogs and cats

Authorised

- Butafosfan
- Cyanocobalamin

Product identification

Medicine name:

Vey Tosal 100 mg/ml + 0.05 mg/ml solution for injection for horses, cattle, dogs and cats

Vey Tosal 100 mg/ml + 0,05 mg/ml Injektionslösung für Rinder, Pferde, Hunde

Active substance:

Butafosfan

Cyanocobalamin

Target species:

Cattle

Horse

Dog

Cat

Route of administration:

Intravenous use

Subcutaneous use

Intramuscular 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:

Germany

Available in:

Germany

Package description:

Glass Vial 1 x 250.0 millilitre(s)

Glass Vial 1 x 100.0 millilitre(s)

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Hybrid application (Article 13(3) of Directive No 2001/82/EC)

Marketing authorisation holder:

Veyx Pharma GmbH

Marketing authorisation date:

30/05/2023

Manufacturing sites for batch release:

Veyx Pharma GmbH

Responsible authority:

Federal Office Of Consumer Protection And Food Safety

Authorisation number:

V7004937.00.00

Date of authorisation status change:

30/05/2023

Reference member state:

Czechia

Procedure number:

CZ/V/0172/001

Concerned member states:

Austria Belgium Bulgaria Croatia Cyprus Denmark Estonia Finland France
Germany Greece Hungary Iceland Ireland Italy Latvia Lithuania
Luxembourg Malta Netherlands Norway Poland Portugal Romania Slovakia
Slovenia Spain Sweden United Kingdom (Northern Ireland)

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

Documents

Combined File of all Documents

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