# Vey Tosal, 100+0,05mg/ml, Solution for injection

Authorised

- Butafosfan
- Cyanocobalamin

# Product identification

### **Medicine name:**

Vey Tosal, 100+0,05mg/ml, Solution for injection Veytosal Vet 100 mg/ml + 0,05 mg/ml injektioneste, liuos

### **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:

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### Cattle

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

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### Horse

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

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# Dog

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Cat

# **Subcutaneous use:**

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# Dog

•

Cat

# Intramuscular use:

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Dog	
• Cat	
Anatomical therapeutic chemical veterinary (ATCvet) codes: QA12CX99	
<b>Legal status of supply:</b> Veterinary medicinal product subject to veterinary prescription	
<b>Authorisation status:</b> Valid	
Authorised in: Finland	
Package description: Glass Vial 1 x 250.0 millilitre(s) Glass Vial 1 x 100.0 millilitre(s)	
Additional information	
Entitlement type: Marketing Authorisation	
<b>Legal basis of product authorisation:</b> Hybrid application (Article 13(3) of Directive No 2001/82/EC)	
<b>Marketing authorisation holder:</b> Veyx Pharma GmbH	
Marketing authorisation date: 5/10/2023	
Manufacturing sites for batch release: Veyx Pharma GmbH	
Responsible authority: Finnish Medicines Agency	

Authorisation number: 39037
Date of authorisation status change: 5/10/2023
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
Summary of Product Characteristics
This document does not exist in this language (English). You can find it in another language below.
Package Leaflet

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