

# Prednisolon ad us. vet 10 mg/ml suspension for injection for cattle, horses, dogs and cats

Authorised

- Prednisolone acetate

## Product identification

### **Medicine name:**

Prednisolon ad us. vet 10 mg/ml suspension for injection for cattle, horses, dogs and cats

### **Active substance:**

Prednisolone acetate

### **Target species:**

Cattle

Dog

Horse

Cat

### **Route of administration:**

Intramuscular use

## Product details

### **Active substance and strength:**

Prednisolone acetate  
10.00 milligram(s) / 1.00 millilitre(s)

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**Pharmaceutical form:**

Suspension for injection

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**Withdrawal period by route of administration:**

**Intramuscular use:**

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**Cattle**

- Meat and offal. 35 day
- Milk. 24 hour

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**Horse**

- Meat and offal. 53 day
- Milk. no withdrawal period

Not authorised for use in lactating mares producing milk for human consumption.

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QH02AB06

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Hungary

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**Package description:**

(ID3) 1200 millilitre(s): unspecified outer container with 12 Vial (glass) each with 100 millilitre(s)

(ID2) 600 millilitre(s): unspecified outer container with 6 Vial (glass) each with 100 millilitre(s)

(ID1) 100 millilitre(s): unspecified outer container with 1 Vial (glass) with 100 millilitre(s)

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Generic application (Article 13(1) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Veyx Pharma GmbH

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**Marketing authorisation date:**

29/10/2018

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**Manufacturing sites for batch release:**

Veyx Pharma GmbH

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**Responsible authority:**

Directorate Of Veterinary Medicinal Products

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**Authorisation number:**

3997/X/18/ NÉBIH ÁTI

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**Date of authorisation status change:**

29/10/2018

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**Reference member state:**

Germany

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**Procedure number:**

DE/V/0162/001

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**Concerned member states:**

Austria Belgium Bulgaria Cyprus Czechia Denmark Estonia France Greece Hungary Iceland Ireland Italy Latvia Lithuania Luxembourg Poland Portugal Romania Slovakia Slovenia Spain United Kingdom (Northern Ireland)

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To consult adverse reactions on veterinary medicinal products please go to [www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Summary of Product Characteristics

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