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)

Pharmaceutical form:
Suspension for injection

Withdrawal period by route of administration:
Intramuscular use:

- **Cattle**
  - Meat and offal. 35 day
  - Milk. 24 hour
- **Horse**
  - Meat and offal. 53 day
  - Milk. no withdrawal period

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QH02AB06

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Greece

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)

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:
Veyx Pharma GmbH

Marketing authorisation date:
2/12/2018

Manufacturing sites for batch release:
Veyx Pharma GmbH

Responsible authority:
National Organization For Medicines

Authorisation number:
71599/04-07-2023/K-0221201

Date of authorisation status change:
3/07/2023

Reference member state:
Germany

Procedure number:
DE/V/0162/001

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)

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

Documents

Summary of Product Characteristics

English (RTF)
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Package Leaflet

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