Ketamidor 100 mg/ml - Injektionslösung für Tiere

Authorised

- Ketamine hydrochloride

Product identification

Medicine name:
Ketamidor 100 mg/ml - Injektionslösung für Tiere
Ketamidor, 100mg/ml, Injekční roztok

Active substance:
Ketamine hydrochloride

Target species:
Dog
Cat
Pig
Cattle
Horse

Route of administration:
Intramuscular use
Intravenous use
Subcutaneous use

Product details

Active substance and strength:

Ketamine hydrochloride
115.33 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:
Intramuscular use:

- **Dog**
- **Cat**
- **Pig**
  - Meat and offal. 0 day

Intravenous use:

- **Cattle**
  - Meat and offal. 0 day
  - Milk. 0 day
- **Dog**
- **Horse**
  - Meat and offal. 0 day
  - Milk. 0 day
- **Cat**

Subcutaneous use:

- **Cat**

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QN01AX03

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

Authorisation status:
Valid

Authorised in:
Czechia

Available in:
Czechia

Package description:
Clear glass vial, type I (Ph. Eur.) with bromobutyl-rubber stopper type I (Ph.Eur.) and aluminium cap, packed in a cardboard box. Package size: 1 x 10 ml
Clear glass vial, type I (Ph. Eur.) with bromobutyl-rubber stopper type I (Ph.Eur.) and aluminium cap, packed in a cardboard box. Package size: 1 x 50 ml
Clear glass vial, type I (Ph. Eur.) with bromobutyl-rubber stopper type I (Ph.Eur.) and aluminium cap, packed in a cardboard box. Package size: 5 x 10 ml

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:
Vetviva Richter GmbH

Marketing authorisation date:
31/08/2016

Manufacturing sites for batch release:
Vetviva Richter GmbH

Responsible authority:
Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:
96/075/16-C

Date of authorisation status change:
31/08/2016

Reference member state:
Austria

Procedure number:
AT/V/0009/001

Concerned member states:
Belgium Czechia Denmark Estonia Finland France Germany Greece Hungary Iceland Ireland Netherlands Poland Portugal 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

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Package Leaflet

This document does not exist in this language (English). You can find it in another language below.

Labelling

This document does not exist in this language (English). You can find it in another language below.

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