ANAESTAMINE 100 MG/ML SOLUTION FOR INJECTION

Authorised

- Ketamine hydrochloride

Product identification

Medicine name:
ANAESTAMINE 100 MG/ML SOLUTION FOR INJECTION

Active substance:
Ketamine hydrochloride

Target species:
Cattle
Pig
Rat
Mouse
Hamster
Guinea pig
Rabbit
Cat
Horse
Horse (mare)
Sheep
Goat
Dog

Route of administration:

Intramuscular use
Intravenous use
Intraperitoneal use

Product details

Active substance and strength:
Ketamine hydrochloride
115.34 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:
Intramuscular use:

- **Cattle**
  - Milk. 0 day
  - Meat and offal. 1 day
- **Pig**
  - Meat and offal. 1 day
- **Horse**
  - Meat and offal. 1 day
- **Horse (mare)**
  - Milk. 0 day
- **Sheep**
  - Milk. 0 day
  - Meat and offal. 1 day

- **Goat**
  - Milk. 0 day
  - Meat and offal. 1 day

Intravenous use:

- **Cattle**
  - Milk. 0 day
  - Meat and offal. 1 day
- **Pig**
  - Meat and offal. 1 day
- **Horse**
  - Meat and offal. 1 day
- **Horse (mare)**
  - Milk. 0 day
- **Sheep**
  - Milk. 0 day
  - Meat and offal. 1 day
- **Goat**
  - Milk. 0 day
  - Meat and offal. 1 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QN01AX03

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Czechia

Package description:
Vial containing 10 ml in a carton box
Vial containing 50 ml in a carton box
Vial containing 25 ml in a carton box

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:
Le Vet. Beheer B.V.

Marketing authorisation date:
25/06/2015

Manufacturing sites for batch release:
Produlab Pharma B.V.

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

Authorisation number:
96/062/15-C

Date of authorisation status change:
25/06/2015

Reference member state:

France

Procedure number:
FR/V/0262/001

Concerned member states:
Austria Belgium Czechia Denmark Estonia Finland Greece Hungary Iceland Ireland Italy Latvia Lithuania Luxembourg Netherlands Norway Poland Portugal Romania Slovakia 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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