KETADORM 100 mg/ml

Authorised

- Ketamine hydrochloride

Product identification

Medicine name:
KETADORM 100 mg/ml

Active substance:
Ketamine hydrochloride

Target species:
Cattle
Sheep
Goat
Horse
Pig
Dog
Cat

Route of administration:
Intravenous use
Intramuscular use
Subcutaneous use

Product details

Active substance and strength:
Ketamine hydrochloride

115.30 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:
Intravenous use:

- **Cattle**
  - Milk. 0 day
  - Meat and offal. 0 day
- **Sheep**
  - Milk. 0 day
  - Meat and offal. 0 day
- **Goat**
  - Milk. 0 day
  - Meat and offal. 0 day
- **Horse**
  - Meat and offal. 0 day

Intramuscular use:

- **Sheep**
  - Milk. 0 day
  - Meat and offal. 0 day
- **Pig**
  - Meat and offal. 0 day
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Dog

- **Cat**

Subcutaneous use:

- **Cat**

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QN01AX03

Legal status of supply:
This information is not available for this product.

Authorisation status:
Valid

Authorised in:
Romania

Package description:
Available only in Romanian
Available only in Romanian
Available only in Romanian

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Bremer Pharma GmbH

Marketing authorisation date:
19/03/2019

Manufacturing sites for batch release:
Bremer Pharma GmbH

Responsible authority:
Institute For Control Of Biological Products And Veterinary Medicines

Authorisation number:
240004

Date of authorisation status change:
4/01/2024

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.
