Ketamidor 10% šķīdums injekcijām liellopiem, zirgiem, aitām, cūkām, kazām, suņiem un kaķiem

Authorised

- Ketamine hydrochloride

Product identification

Medicine name:
Ketamidor 10% šķīdums injekcijām liellopiem, zirgiem, aitām, cūkām, kazām, suņiem un kaķiem

Active substance:
Ketamine hydrochloride

Target species:
Horse
Cattle
Sheep
Goat
Dog
Cat
Pig

Route of administration:
Intravenous use

Intramuscular use
Subcutaneous use

Product details

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

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:
Intravenous use:

- **Horse**
  - Not specified. 0 day
- **Cattle**
  - Not specified. 0 day
- **Sheep**
  - Not specified. 0 day
- **Goat**
  - Not specified. 0 day
- **Pig**
  - Not specified. 0 day

Intramuscular use:

- **Sheep**

- Not specified. 0 day

- **Goat**
  - Not specified. 0 day

- **Pig**
  - Not specified. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QN01AX03

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

Authorisation status:
Valid

Authorised in:
Latvia

Available in:
Latvia

Package description:
Available only in Latvian

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Well-established use application (Article 13a of Directive No 2001/82/EC)

Marketing authorisation holder:
Vetviva Richter GmbH

Marketing authorisation date:
16/02/2001

Manufacturing sites for batch release:
Richter Pharma AG

Responsible authority:
Food And Veterinary Service

Authorisation number:
V/NRP/01/1290

Date of authorisation status change:
18/02/2001

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

Documents

Combined File of all Documents

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