File downloaded on 2026-11-09
Source URL: https://medicines.health.europa.eu/veterinary/en/600000073113

Ketamin 100 mg/ml

Authorised

- Ketamine hydrochloride

Product identification

Medicine name:
Ketamin 100 mg/ml

Active substance:
Ketamine hydrochloride

Target species:
Cattle
Pig
Dog
Cat
Sheep
Goat
Horse

Route of administration:
Intramuscular use
Subcutaneous use
Intravenous 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**
  - Meat and offal. 3 day
  - Milk. 0 day
- **Pig**
  - Meat and offal. 3 day
- **Sheep**
  - Meat and offal. 3 day
  - Milk. 0 day
- **Goat**
  - Meat and offal. 3 day
  - Milk. 0 day

Intravenous use:

- **Horse**
  - Meat and offal. 3 day
  - Milk. no withdrawal period

Nicht bei Stuten anwenden, deren Milch für den menschlichen Verzehr vorgesehen ist.

-

Cattle

- Meat and offal. 3 day
- Milk. 0 day

•

Pig

- Meat and offal. 3 day

•

Sheep

- Meat and offal. 3 day
- Milk. 0 day

•

Goat

- Meat and offal. 3 day
- Milk. 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:
Germany

Available in:
Germany

Package description:
Available only in German
Available only in German
Available only in German
Available only in German

Available only in German
Available only in German
Available only in German
Available only in German
Available only in German

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:
CP-Pharma Handelsgesellschaft mbH

Marketing authorisation date:
21/05/2013

Manufacturing sites for batch release:
CP-Pharma Handelsgesellschaft mbH

Responsible authority:
Federal Office Of Consumer Protection And Food Safety

Authorisation number:
401650.00.00

Date of authorisation status change:
14/11/2019

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

Documents

Combined File of all Documents

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

2401650-parde-20231002.pdf