

## KETAMINOL 10

Not authorised

- Ketamine hydrochloride

## Product identification

**Medicine name:**

KETAMINOL 10

**Active substance:**

Ketamine hydrochloride

**Target species:**

Horse

Cattle (cow)

Cattle (calf)

Pig

Dog

Cat

Sheep

Goat

Guinea pig

Hamster

Rabbit

Chicken

Bear

**Route of administration:**

Intravenous use

Intramuscular 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:**

#### **Intravenous use:**

- 

#### **Horse**

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

- 

#### **Cattle (cow)**

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

- 

#### **Cattle (calf)**

- Meat and offal. 0 day

- 

#### **Pig**

- Meat and offal. 0 day

- 

#### **Sheep**

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

-

**Goat**

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

**Intramuscular use:**

•

**Pig**

- Meat and offal. 0 day

•

**Sheep**

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

•

**Goat**

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

•

**Rabbit**

- Meat and offal. 0 day

•

**Chicken**

- Meat. 0 day

•

**Bear**

- Meat. 0 day

---

**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QN01AX03

---

**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

---

**Authorisation status:**

Surrendered

---

**Authorised in:**

Romania

---

**Package description:**

Available only in Romanian

---

## Additional information

**Entitlement type:**

Marketing Authorisation

---

**Legal basis of product authorisation:**

Complete application (stand-alone) - Directive No 2001/82/EC

---

**Marketing authorisation holder:**

Intervet International B.V.

---

**Marketing authorisation date:**

22/06/2005

---

**Manufacturing sites for batch release:**

Intervet Deutschland GmbH

---

**Responsible authority:**

Institute For Control Of Biological Products And Veterinary Medicines

---

**Authorisation number:**

150193

---

**Date of authorisation status change:**

17/03/2025

---

To consult adverse reactions on veterinary medicinal products please go to  
[www.adrreports.eu/vet](http://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.