

# Ketamidor 100 mg/ml - Injektionslösung für Tiere

Authorised

- Ketamine hydrochloride

## Product identification

**Medicine name:**

Ketamidor 100 mg/ml - Injektionslösung für Tiere

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**Active substance:**

Ketamine hydrochloride

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**Target species:**

Dog

Cat

Pig

Cattle

Horse

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**Route of administration:**

Intramuscular use

Intravenous use

Subcutaneous use

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## Product details

**Active substance and strength:**

Ketamine hydrochloride  
115.33 milligram(s) / 1.00 millilitre(s)

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**Pharmaceutical form:**

Solution for injection

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**Withdrawal period by route of administration:**

**Intramuscular use:**

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**Pig**

- Meat and offal. 0 day

**Intravenous use:**

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**Cattle**

- Meat and offal. 0 day

- Milk. 0 day

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**Horse**

- Meat and offal. 0 day

- Milk. 0 day

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

QN01AX03

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Finland

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**Available in:**

Finland

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**Package description:**

Clear glass vial, type I (Ph. Eur.) with bromobutyl-rubber stopper type I (Ph.Eur.) and aluminium cap, packed in a cardboard box. Package size: 1 x 10 ml

Clear glass vial, type I (Ph. Eur.) with bromobutyl-rubber stopper type I (Ph.Eur.) and aluminium cap, packed in a cardboard box. Package size: 1 x 50 ml

Clear glass vial, type I (Ph. Eur.) with bromobutyl-rubber stopper type I (Ph.Eur.) and aluminium cap, packed in a cardboard box. Package size: 5 x 10 ml

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Generic application (Article 13(1) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Vetviva Richter GmbH

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**Marketing authorisation date:**

16/01/2013

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**Manufacturing sites for batch release:**

Vetviva Richter GmbH

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**Responsible authority:**

Finnish Medicines Agency

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**Authorisation number:**

30400

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**Date of authorisation status change:**

16/01/2013

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**Reference member state:**

Austria

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**Procedure number:**

AT/V/0009/001

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**Concerned member states:**

Belgium Czechia Denmark Estonia Finland France Germany Greece  
Hungary Iceland Ireland Netherlands Poland Portugal Slovenia Spain  
Sweden United Kingdom (Northern Ireland)

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

## Documents

### Summary of Product Characteristics

English (PDF)

Published on: 23/04/2025

Updated on: 13/03/2026

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### Package Leaflet

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

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