

# ANAESTAMINE 100 MG/ML SOLUTION FOR INJECTION

Authorised

- Ketamine hydrochloride

## Product identification

**Medicine name:**

ANAESTAMINE 100 MG/ML SOLUTION FOR INJECTION

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**Active substance:**

Ketamine hydrochloride

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**Target species:**

Cattle

Pig

Rat

Mouse

Hamster

Guinea pig

Rabbit

Cat

Horse

Horse (mare)

Sheep

Goat

Dog

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**Route of administration:**

Intramuscular use  
Intravenous use  
Intraperitoneal use

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## Product details

### **Active substance and strength:**

Ketamine hydrochloride  
115.34 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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#### **Cattle**

- Milk. 0 day
- Meat and offal. 1 day

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#### **Pig**

- Meat and offal. 1 day

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#### **Horse**

- Meat and offal. 1 day

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#### **Horse (mare)**

- Milk. 0 day

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#### **Sheep**

- Milk. 0 day
- Meat and offal. 1 day

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**Goat**

- Milk. 0 day
- Meat and offal. 1 day

**Intravenous use:**

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**Cattle**

- Milk. 0 day
- Meat and offal. 1 day

- 

**Pig**

- Meat and offal. 1 day

- 

**Horse**

- Meat and offal. 1 day

- 

**Horse (mare)**

- Milk. 0 day

- 

**Sheep**

- Milk. 0 day
- Meat and offal. 1 day

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**Goat**

- Milk. 0 day
- Meat and offal. 1 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:**

Romania

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**Package description:**

Vial containing 10 ml in a carton box

Vial containing 50 ml in a carton box

Vial containing 25 ml in a carton box

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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:**

Le Vet. Beheer B.V.

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**Marketing authorisation date:**

18/02/2015

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**Manufacturing sites for batch release:**

Produlab Pharma B.V.

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**Responsible authority:**

Institute For Control Of Biological Products And Veterinary Medicines

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**Authorisation number:**

190183

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**Date of authorisation status change:**

4/11/2019

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**Reference member state:**

France

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**Procedure number:**

FR/V/0262/001

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**Concerned member states:**

Austria Belgium Czechia Denmark Estonia Finland Greece Hungary Iceland  
Ireland Italy Latvia Lithuania Luxembourg Netherlands Norway Poland  
Portugal Romania Slovakia 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

Combined File of all Documents

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