

KETADORM 100 mg/ml

Authorised

- Ketamine hydrochloride

Product identification

Medicine name:

KETADORM 100 mg/ml

Active substance:

Ketamine hydrochloride

Target species:

Cattle

Sheep

Goat

Horse

Pig

Dog

Cat

Route of administration:

Intravenous use

Intramuscular use

Subcutaneous use

Product details

Active substance and strength:

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

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Intravenous use:

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Cattle

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

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Sheep

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

•

Goat

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

•

Horse

- Meat and offal. 0 day

Intramuscular use:

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Sheep

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

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Pig

- Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QN01AX03

Legal status of supply:

This information is not available for this product.

Authorisation status:

Valid

Authorised in:

Romania

Package description:

Available only in Romanian

Available only in Romanian

Available only in Romanian

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:

Bremer Pharma GmbH

Marketing authorisation date:

19/03/2019

Manufacturing sites for batch release:

Laboratorios Karizoo S.A.

Responsible authority:

Institute For Control Of Biological Products And Veterinary Medicines

Authorisation number:

240004

Date of authorisation status change:

16/11/2025

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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