

NARKAMON 50 mg/ml injekčný roztok

Authorised

- Ketamine hydrochloride

Product identification

Medicine name:

NARKAMON 50 mg/ml injekčný roztok

Active substance:

Ketamine hydrochloride

Target species:

Cattle (calf)

Sheep

Goat

Dog

Cat

Guinea pig

Monkey

Parrot

Reptile

Antelope

Roe deer

Pigeon

Red deer

Mouse

Rat
Horse

Route of administration:

Intramuscular use
Intraperitoneal use
Intravenous use

Product details

Active substance and strength:

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

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Intramuscular use:

•

Cattle (calf)

- Meat and offal. 24 hour

•

Sheep

- Meat and offal. 24 hour

- Milk. 0 day
milk zero days

•

Goat

- Meat and offal. 24 hour

- Milk. 0 day
milk zero days

•

Dog

- Not applicable. 0 day Not applicable

•

Cat

- Not applicable. 0 day Not applicable

•

Guinea pig

- Not applicable. 0 day Not applicable

•

Monkey

- Not applicable. 0 day Not applicable

•

Parrot

- Not applicable. 0 day Not applicable

•

Reptile

- Not applicable. 0 day Not applicable

•

Antelope

- Meat and offal. 24 hour

•

Roe deer

- Meat and offal. 24 hour

•

Pigeon

- Meat and offal. 24 hour

•

Red deer

- Meat and offal. 24 hour

Intraperitoneal use:

•

Mouse

- Not applicable. 0 day Not applicable

•

Rat

- Not applicable. 0 day Not applicable

Intravenous use:

•

Horse

- Meat and offal. 24 hour
- Milk. 0 day milkmzreo days

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Dog

- Not applicable. 0 day Not applicable

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QN01AX03

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Slovakia

Package description:

Available only in Slovak

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:

Bioveta a.s.

Marketing authorisation date:

26/09/1988

Manufacturing sites for batch release:

Bioveta a.s.

Responsible authority:

Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:

99/172/88-C/S

Date of authorisation status change:

26/09/1988

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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