NARKAMON 100 MG/ML SOLUTION INJECTABLE

Authorised

- Ketamine hydrochloride

Product identification

Medicine name:
NARKAMON 100 MG/ML SOLUTION INJECTABLE

Active substance:
Ketamine hydrochloride

Target species:
Dog
Cat
Donkey
Horse

Route of administration:
Intramuscular use
Intravenous 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:

- **Donkey**
  - Meat and offal. 1 day
  - Milk. 24 hour
- **Horse**
  - Meat and offal. 1 day
  - Milk. 24 hour

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QN01AX03

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
France

Package description:
Available only in French
Available only in French

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:

Hybrid application (Article 13(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Osalia

Marketing authorisation date:
20/07/2022

Manufacturing sites for batch release:
Bioveta a.s.

Responsible authority:
French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:
FR/V/7340298 3/2022

Date of authorisation status change:
20/07/2022

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Summary of Product Characteristics

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

Package Leaflet and Labelling

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

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