Anesketin 100 mg/ml Solution for Injection

Authorised

- Ketamine hydrochloride

Product identification

Medicine name:
Anesketin 100 mg/ml Solution for Injection
ANESKETIN 100 MG/ML SOLUTION INJECTABLE

Active substance:
Ketamine hydrochloride

Target species:
Horse
Dog
Cat

Route of administration:
Intramuscular use
Intravenous use
Subcutaneous use

Product details

Active substance and strength:
Ketamine hydrochloride
115.40 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Horse**
  - Milk. 1 day
  - Meat and offal. 1 day
- **Dog**
- **Cat**

Intravenous use:

- **Horse**
  - Milk. 1 day
  - Meat and offal. 1 day
- **Dog**
- **Cat**

Subcutaneous use:

- **Horse**
  - Milk. 1 day
  - Meat and offal. 1 day
- **Dog**
- **Cat**

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QN01AX03

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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Authorisation status:

Valid

Authorised in:

France

Package description:

Clear colourless type I glass vials with bromobutyl rubber stoppers and aluminium caps filled with 50 ml, 1 vial in a cardboard box
Clear colourless type I glass vials with bromobutyl rubber stoppers and aluminium caps filled with 5 ml, 1 vial in a cardboard box
Clear colourless type I glass vials with bromobutyl rubber stoppers and aluminium caps filled with 30 ml, 1 vial in a cardboard box
Clear colourless type I glass vials with bromobutyl rubber stoppers and aluminium caps filled with 25 ml, 1 vial in a cardboard box
Clear colourless type I glass vials with bromobutyl rubber stoppers and aluminium caps filled with 20 ml, 1 vial in a cardboard box
Clear colourless type I glass vials with bromobutyl rubber stoppers and aluminium caps filled with 10 ml, 1 vial in a cardboard box

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:
Eurovet Animal Health B.V.

Marketing authorisation date:
This information is not available for this product.

Manufacturing sites for batch release:
Eurovet Animal Health B.V.

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

Authorisation number:
FR/V/9465271 4/2013

Date of authorisation status change:
17/05/2018

Reference member state:
Netherlands

Procedure number:
NL/V/0278/001

Concerned member states:
Belgium Denmark France Germany Spain United Kingdom (Northern Ireland)

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.

Combined File of all Documents

Package Leaflet and Labelling

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