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Anesketin 100 mg/ml Solution for Injection for Dogs, Cats and Horses

Authorised

- Ketamine hydrochloride

Product identification

Medicine name:
Anesketin 100 mg/ml Solution for Injection for Dogs, Cats and Horses

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

Intravenous use:

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

Subcutaneous use:

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QN01AX03

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

Authorisation status:
Valid

Authorised in:
United Kingdom (Northern Ireland)

Available in:
United Kingdom (Northern Ireland)

Package description:
Clear colourless type I glass vials with bromobutyl rubber stoppers and aluminium caps filled with 10 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 25 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 5 ml, 1 vial in a cardboard box
Clear colourless type I glass vials with bromobutyl rubber stoppers and aluminium caps filled with 50 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:
10/07/2013

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

Responsible authority:
The Veterinary Medicines Directorate

Authorisation number:

Vm 16849/3002

Date of authorisation status change:
18/05/2024

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

Combined File of all Documents