Ketavet 100 mg/ml solution for injection for dogs, cats and horses

Authorised

- Ketamine

Product identification

Medicine name:
Ketavet 100 mg/ml solution for injection for dogs, cats and horses

Active substance:
Ketamine

Target species:
Dog
Cat
Horse

Route of administration:
Intramuscular use
Intravenous use
Subcutaneous use

Product details

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

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:

Intravenous use:

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

Package description:
Clear colourless type I glass vials with chlorobutyl rubber stoppers and aluminium flip off seals. Cardboard box containing 1 vial of 50 ml.
Clear colourless type I glass vials with chlorobutyl rubber stoppers and aluminium flip off seals. Cardboard box containing 1 vial of 10 ml.

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:

Zoetis Belgium S.A.

Marketing authorisation date:
20/03/2015

Manufacturing sites for batch release:
Zoetis Manufacturing & Research Spain S.L.

Responsible authority:
Health Products Regulatory Authority

Authorisation number:
VPA10387/036/001

Date of authorisation status change:
20/03/2015

Reference member state:
Ireland

Procedure number:
IE/V/0646/001

Concerned member states:
Austria Cyprus Germany Greece Italy Portugal 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

English (PDF)

Published on: 11/04/2023

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Package Leaflet

Labelling

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