Genabil 100 mg/ml raztopina za injiciranje za govedo, prašiče, konje, ovce in pse

Authorised

- Menbutone

Product identification

Medicine name:
Genabil 100 mg/ml raztopina za injiciranje za govedo, prašiče, konje, ovce in pse

Active substance:
Menbutone

Target species:
Cattle
Horse
Pig
Sheep
Dog

Route of administration:
Intravenous use
Intramuscular use

Product details

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

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:
Intravenous use:

- **Cattle**
  - Meat and offal. 3 day Meso in organi: 3 dni.
  - Milk. 2 day Mleko: 2 dneva.
- **Horse**
  - Meat and offal. 3 day Meso in organi: 3 dni.
  - Milk. 2 day

Mleko: 2 dneva. Ne uporabite pri kobilah, katerih mleko je namenjeno prehrani ljudi.
Not for use in mares whose milk is intended for human consumption.

- **Pig**
  - Meat and offal. 3 day Meso in organi: 3 dni.
- **Sheep**
  - Meat and offal. 3 day Meso in organi: 3 dni.
  - Milk. 2 day Mleko: 2 dneva.

Intramuscular use:

- **Cattle**
  - Meat and offal. 3 day — Meso in organi: 3 dni.
  - Milk. 2 day — Mleko: 2 dneva.
- **Pig**
  - Meat and offal. 3 day — Meso in organi: 3 dni.
- **Sheep**
  - Milk. 2 day — Mleko: 2 dneva.

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QA05AX90

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

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Authorisation status:
Valid

Authorised in:
Slovenia

Package description:
Available only in Slovenian

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:

Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:
Boehringer Ingelheim Vetmedica GmbH

Marketing authorisation date:
19/11/2021

Manufacturing sites for batch release:
Labiana Life Sciences S.A.

Responsible authority:
Agency For Medicinal Products And Medical Devices Of The Republic Of Slovenia

Authorisation number:
NP/V/0152/001

Date of authorisation status change:
14/08/2002

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

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

Labelling

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