VETBUTON 100 MG/ML SOLUTION FOR INJECTION FOR CATTLE, PIGS, HORSES, SHEEP AND GOATS

Authorised

- Menbutone

Product identification

Medicine name:
VETBUTON 100 MG/ML SOLUTION FOR INJECTION FOR CATTLE, PIGS, HORSES, SHEEP AND GOATS

Active substance:
Menbutone

Target species:
Goat
Pig
Sheep
Cattle
Horse

Route of administration:
Intramuscular use
Intravenous 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:
Intramuscular use:

- **Goat**
  - Meat and offal. 0 day
  - Milk. 0 day
- **Pig**
  - Meat and offal. 0 day
- **Sheep**
  - Meat and offal. 0 day
  - Milk. 0 day

Intravenous use:

- **Cattle**
  - Meat and offal. 0 day
  - Milk. 0 day
- **Pig**
  - Meat and offal. 0 day
-

Horse

- Meat and offal. 0 day
- Milk. 0 day

•

Sheep

- Meat and offal. 0 day
- Milk. 0 day

•

Goat

- Meat and offal. 0 day
- Milk. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QA05AX90

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

Authorisation status:
Valid

Authorised in:
Netherlands

Available in:
Netherlands

Package description:
Cardboard box with one 100 ml natural multi-layer (COEX) PP/EVOH/PP vial closed with bromobutyl rubber stopper and aluminium and plastic flip capsule.

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:
Vet-Agro Multi-Trade Company Sp. z o.o.

Marketing authorisation date:
3/08/2019

Manufacturing sites for batch release:
Przedsiebiorstwo Wielobranzowe Vet-Agro Sp. z o.o.

Responsible authority:
Medicines Evaluation Board

Authorisation number:
REG NL 123496

Date of authorisation status change:
29/06/2022

Reference member state:
France

Procedure number:
FR/V/0324/001

Concerned member states:
Belgium Croatia Netherlands Portugal Romania Slovenia Spain Sweden

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

Documents

Combined File of all Documents

English (PDF)
Published on: 14/03/2026
Download

eu-puar-frv0324001-mr-rpe522-en.pdf