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:
Belgium

Available in:
Belgium

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:
Przedsiebiorstwo Wielobranzowe Vet-Agro Sp. z o.o.

Marketing authorisation date:
22/08/2019

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

Responsible authority:
Federal Agency For Medicines And Health Products

Authorisation number:
BE-V544782

Date of authorisation status change:
22/08/2019

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

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.

Combined File of all Documents

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