

# 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)

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### **Pharmaceutical form:**

Solution for injection

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### **Withdrawal period by route of administration:**

#### **Intramuscular use:**

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##### **Goat**

- Meat and offal. 0 day

- Milk. 0 day

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##### **Pig**

- Meat and offal. 0 day

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##### **Sheep**

- Meat and offal. 0 day

- Milk. 0 day

#### **Intravenous use:**

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##### **Cattle**

- Meat and offal. 0 day

- Milk. 0 day

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##### **Pig**

- Meat and offal. 0 day

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**Horse**

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

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**Sheep**

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

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**Goat**

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

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QA05AX90

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Croatia

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**Available in:**

Croatia

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**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.

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Generic application (Article 13(1) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

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

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**Marketing authorisation date:**

9/08/2019

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**Manufacturing sites for batch release:**

Przedsiębiorstwo Wielobranzowe Vet-Agro Sp. z o.o.

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**Responsible authority:**

Ministry Of Agriculture Veterinary And Food Safety Directorate

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**Authorisation number:**

UP/I-322-05/19-01/468

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**Date of authorisation status change:**

9/08/2019

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**Reference member state:**

France

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**Procedure number:**

FR/V/0324/001

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**Concerned member states:**

Belgium Croatia Netherlands Portugal Romania Slovenia Spain Sweden

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To consult adverse reactions on veterinary medicinal products please go to  
[www.adrreports.eu/vet](http://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.

Combined File of all Documents

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