# Virbamec 1% 10 mg/ml Solution injectable

Authorised

- Ivermectin

## Product identification

**Medicine name:**
Virbamec 1% 10 mg/ml Solution injectable
Virbamec 1% 10 mg/ml Oplossing voor injectie
Virbamec 1% 10 mg/ml Injektionslösung

**Active substance:**
Ivermectin

**Target species:**
Sheep
Pig
Cattle

**Route of administration:**
Subcutaneous use

## Product details

**Active substance and strength:**
Ivermectin
10.00 milligram(s) / 1.00 millilitre(s)

**Pharmaceutical form:**

Solution for injection

**Withdrawal period by route of administration:**

**Subcutaneous use:**

- **Sheep**

  - Meat and offal. 45 day

  - Milk. no withdrawal period

Do not use in animals producing milk for human consumption

  - Milk. no withdrawal period

Do not use in sheep which are intended to produce milk for human consumption within 60 days of lambing

- **Pig**

  - Meat and offal. 35 day

- **Cattle**

  - Meat and offal. 49 day

  - Milk. no withdrawal period

Do not use in animals producing milk for human consumption

  - Milk. no withdrawal period

Do not use in non-lactating dairy cows, including pregnant dairy heifers, within 60 days of calving

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

QP54AA01

**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

**Authorisation status:**

Valid

**Authorised in:**

Belgium

**Package description:**

Virbamec 1% 10 mg/ml Solution for injection Vial of 1 l

Virbamec 1% 10 mg/ml Solution for injection Vial of 500 ml
Virbamec 1% 10 mg/ml Solution for injection Vial of 200 ml

## Additional information

**Entitlement type:**
Marketing Authorisation

**Legal basis of product authorisation:**
Generic application (Article 18 of Regulation (EU) 2019/6)

**Marketing authorisation holder:**
Virbac

**Marketing authorisation date:**
This information is not available for this product.

**Manufacturing sites for batch release:**
VIRBAC
Sofarimex-Industria Quimica E Farmaceutica S.A.

**Responsible authority:**
Federal Agency For Medicines And Health Products

**Authorisation number:**
BE-V277216

**Date of authorisation status change:**
23/07/2018

**Generic of:**
600000086014

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

## Documents

Package Leaflet

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

Summary of Product Characteristics

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.

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