# Intermectin Injection 10 mg/ml solution for injection for cattle, sheep and pigs

Authorised

- Ivermectin

## Product identification

**Medicine name:**
Intermectin Injection 10 mg/ml solution for injection for cattle, sheep and pigs
Intermectin 10 mg/ml raztopina za injiciranje za govedo, ovce in prašiče

**Active substance:**
Ivermectin

**Target species:**
Cattle
Sheep
Pig

**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:**

- **Cattle**
  - Meat and offal. 49 day
- **Sheep**
  - Meat and offal. 63 day
- **Pig**
  - Meat and offal. 28 day

**Anatomical therapeutic chemical veterinary (ATCvet) codes:**
QP54AA01

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

**Authorisation status:**
Valid

**Authorised in:**
Slovenia

**Package description:**
Clear glass bottles (type II) closed with a bromobutyl rubber stopper and sealed with an aluminium cap packed into an outer cardboard box
Clear glass bottles (type II) closed with a bromobutyl rubber stopper and sealed with an aluminium flip-off cap with polypropylene cover packed into an outer cardboard box

## Additional information

**Entitlement type:**
Marketing Authorisation

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

**Marketing authorisation holder:**
Interchemie Werken De Adelaar B.V.

**Marketing authorisation date:**
7/04/2025

**Manufacturing sites for batch release:**
Interchemie Werken De Adelaar Eesti AS

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

**Authorisation number:**
MR/V/0817/001

**Date of authorisation status change:**
7/04/2025

**Reference member state:**
Netherlands

**Procedure number:**
NL/V/0431/001

**Concerned member states:**
Austria Croatia Cyprus Czechia France Germany Greece Hungary Italy Latvia Lithuania Malta Portugal Slovakia Slovenia Spain

**Generic of:**
600000065934

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.