# LV Intermectin 10 mg/ml šķīdums injekcijām liellopiem, aitām, kazām un cūkām

Authorised

- Ivermectin

## Product identification

**Medicine name:**
LV Intermectin 10 mg/ml šķīdums injekcijām liellopiem, aitām, kazām un cūkām

**Active substance:**
Ivermectin

**Target species:**
Sheep
Pig
Cattle
Goat

**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. 49 day
  - Milk. no withdrawal period

Nav reģistrēts lietošanai dzīvniekiem, kuru pienu paredzēts izmantot cilvēku uzturā.

- **Pig**
  - Meat and offal. 28 day

- **Cattle**
  - Meat and offal. 49 day
  - Milk. no withdrawal period

Nav reģistrēts lietošanai dzīvniekiem, kuru pienu paredzēts izmantot cilvēku uzturā.

- **Goat**
  - Milk. no withdrawal period

Nav reģistrēts lietošanai dzīvniekiem, kuru pienu paredzēts izmantot cilvēku uzturā.

  - Meat and offal. 49 day

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

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

**Authorisation status:**

Valid

**Authorised in:**
Latvia

**Package description:**
Available only in Latvian
Available only in Latvian
Available only in Latvian
Available only in Latvian
Available only in Latvian
Available only in Latvian
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Available only in Latvian

## Additional information

**Entitlement type:**
Marketing Authorisation

**Legal basis of product authorisation:**
Well-established use application (Article 13a of Directive No 2001/82/EC)

**Marketing authorisation holder:**
Interchemie Werken De Adelaar Eesti AS

**Marketing authorisation date:**
28/05/2012

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

**Responsible authority:**
Food And Veterinary Service

**Authorisation number:**
V/NRP/12/0041

**Date of authorisation status change:**

28/05/2012

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

## Documents

Combined File of all Documents

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