

# CEVAMEC 1 %, injekcinis tirpalas

Not  
authorised

- Ivermectin

## Product identification

**Medicine name:**

CEVAMEC 1 %, injekcinis tirpalas

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**Active substance:**

Ivermectin

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**Target species:**

Cattle

Sheep

Goat

Pig

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**Route of administration:**

Subcutaneous use

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## Product details

**Active substance and strength:**

Ivermectin

10.00 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:****Subcutaneous use:**

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**Cattle**

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

Not authorized for use in lactating females, whose milk will be used for human consumption and pregnant females, whose milk will be used for human consumption, 28 days before delivery.

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**Sheep**

- Meat and offal. 21 day
- Milk. no withdrawal period

Not authorized for use in lactating females, whose milk will be used for human consumption and pregnant females, whose milk will be used for human consumption, 28 days before delivery.

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**Goat**

- Meat and offal. 28 day
- Milk. no withdrawal period

Not authorized for use in lactating females, whose milk will be used for human consumption and pregnant females, whose milk will be used for human consumption, 28 days before delivery.

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**Pig**

- Meat and offal. 28 day

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

QP54AA01

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription except for some pack sizes

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**Authorisation status:**

Surrendered

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**Authorised in:**

Lithuania

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**Package description:**

Available only in [Lithuanian](#)

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Legal basis not covered by Directive 2001/82/EC

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**Marketing authorisation holder:**

Ceva Sante Animale

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**Marketing authorisation date:**

23/03/2000

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**Manufacturing sites for batch release:**

Ceva-Phylaxia Zrt.

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**Responsible authority:**

State Food And Veterinary Service

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**Authorisation number:**

LT/2/00/1080/001-005

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**Date of authorisation status change:**

6/11/2010

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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

RV1080.pdf