

INTERMEC, 10 mg/ml injekcinis tirpalas galvijams, avims, ožkoms ir kiaulėms

Authorised

- Ivermectin

Product identification

Medicine name:

INTERMEC, 10 mg/ml injekcinis tirpalas galvijams, avims, ožkoms ir kiaulėms

Active substance:

Ivermectin

Target species:

Cattle

Goat

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

Not authorized for use in females producing milk for human consumption.

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Goat

- Meat and offal. 49 day

Not authorized for use in females producing milk for human consumption.

-

Sheep

- Meat and offal. 49 day

Not authorized for use in females producing milk for human consumption.

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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:

Lithuania

Package description:

Available only in [Lithuanian](#)

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Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

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

Marketing authorisation holder:

Interchemie Werken De Adelaar B.V.

Marketing authorisation date:

9/12/2010

Manufacturing sites for batch release:

Interchemie Werken De Adelaar Eesti AS

PharmaPark Production OÜ

Responsible authority:

State Food And Veterinary Service

Authorisation number:

LT/2/10/2009/001-006

Date of authorisation status change:

24/09/2026

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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