

# Noromectin 10 mg/ml injekčný roztok

Authorised

- Ivermectin

## Product identification

**Medicine name:**

Noromectin 10 mg/ml injekčný roztok

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**Active substance:**

Ivermectin

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**Target species:**

Cattle

Sheep

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

Do not use in cows whose milk is intended for human consumption.

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**Sheep**

- Meat and offal. 42 day
- Milk. no withdrawal period

Do not use in sheep whose milk is intended for human consumption.

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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

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**Authorisation status:**

Valid

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**Authorised in:**

Slovakia

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**Available in:**

Slovakia

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**Package description:**

Available only in Slovak

Available only in [Slovak](#)

Available only in [Slovak](#)

Available only in [Slovak](#)

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Norbrook Laboratories (Ireland) Limited

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**Marketing authorisation date:**

29/11/2002

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**Manufacturing sites for batch release:**

Norbrook Manufacturing Limited

Norbrook Laboratories Limited

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**Responsible authority:**

Institute For State Control Of Veterinary Biologicals And Medicaments

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**Authorisation number:**

96/084/02-S

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**Date of authorisation status change:**

29/11/2002

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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

Combined File of all Documents

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