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# Taurador 10 mg/ml Solution for Injection for cattle, sheep & pigs

Authorised

- Doramectin

## Product identification

**Medicine name:**
Taurador 10 mg/ml Solution for Injection for cattle, sheep & pigs

**Active substance:**
Doramectin

**Target species:**
Cattle
Sheep
Pig

**Route of administration:**
Subcutaneous use
Intramuscular use

## Product details

**Active substance and strength:**
Doramectin
10.00 milligram(s) / 1.00 millilitre(s)

**Pharmaceutical form:**
Solution for injection

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**Withdrawal period by route of administration:**
**Subcutaneous use:**

- **Cattle**
  - Meat and offal. 70 day
  - Milk. no withdrawal period

Not authorised for use in animals producing milk for human consumption. Do not use in pregnant cows or heifers which are intended to produce milk for human consumption within 2 months of expected parturition.

**Intramuscular use:**

- **Sheep**
  - Meat and offal. 70 day
  - Milk. no withdrawal period

Not authorised for use in animals producing milk for human consumption. Do not use in pregnant ewes which are intended to produce milk for human consumption within 70 days of expected parturition.

- **Pig**
  - Meat and offal. 77 day

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**
QP54AA03

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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:**

Romania

**Package description:**
(ID1) 100 millilitre(s): container (Plastic) with 1 Vial (Glass) with 100 millilitre(s), closed with Stopper (Rubber) and Lid (Aluminium)
(ID2) 250 millilitre(s): container (Plastic) with 1 Vial (Glass) with 250 millilitre(s), closed with Lid (Aluminium) and Stopper (Rubber)
(ID3) 500 millilitre(s): container (Plastic) with 1 Vial (Glass) with 500 millilitre(s), closed with Stopper (Rubber) and Lid (Aluminium)

## Additional information

**Entitlement type:**
Marketing Authorisation

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

**Marketing authorisation holder:**
Norbrook Laboratories (Ireland) Limited

**Marketing authorisation date:**
20/01/2025

**Manufacturing sites for batch release:**
Norbrook Laboratories Limited
Norbrook Manufacturing Limited

**Responsible authority:**
Institute For Control Of Biological Products And Veterinary Medicines

**Authorisation number:**
250014

**Date of authorisation status change:**
4/03/2026

**Reference member state:**
Germany

**Procedure number:**
DE/V/0345/001

**Concerned member states:**
Belgium Czechia France Hungary Ireland Netherlands Portugal Romania Slovakia Spain

**Generic of:**
600000073029
600000060573
600000085745

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## Documents

Combined File of all Documents

English (PDF)
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