File downloaded on 2026-11-09
Source URL: https://medicines.health.europa.eu/veterinary/en/700000140594

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

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QP54AA03

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

---

Authorisation status:
Valid

Authorised in:

Czechia

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:
18/02/2025

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

Responsible authority:
Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:
96/007/25-C

Date of authorisation status change:
18/02/2025

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

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

Documents

Summary of Product Characteristics

This document does not exist in this language (English). You can find it in another language below.

Package Leaflet

This document does not exist in this language (English). You can find it in another language below.

Labelling

This document does not exist in this language (English). You can find it in another language below.

Combined File of all Documents

7015912-paren-20250130.pdf