

# Doramax 10 mg/ml Solution for Injection for Cattle, Sheep and Pigs

Authorised

- Doramectin

## Product identification

**Medicine name:**

Doramax 10 mg/ml Solution for Injection for Cattle, Sheep and Pigs

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**Active substance:**

Doramectin

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**Target species:**

Sheep  
Pig  
Cattle

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**Route of administration:**

Intramuscular use  
Subcutaneous use

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## Product details

**Active substance and strength:**

Doramectin

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:**

**Intramuscular use:**

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**Sheep**

- Meat and offal. 70 day

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**Pig**

- Meat and offal. 77 day

**Subcutaneous use:**

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**Cattle**

- Meat and offal. 70 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:**

Germany

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**Package description:**

Type II amber glass multi-dose vials, with chlorobutyl rubber stoppers secured with an aluminium cap. Package sizes: Cardboard box or Plastic sleeve containing 1 vial of 500 ml.

Type II amber glass multi-dose vials, with chlorobutyl rubber stoppers secured with an aluminium cap. Package sizes: Cardboard box or Plastic sleeve containing 1 vial of

250 ml.

Type II amber glass multi-dose vials, with chlorobutyl rubber stoppers secured with an aluminium cap. Package sizes: Cardboard box or Plastic sleeve containing 1 vial of 50 ml.

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

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

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**Marketing authorisation holder:**

Chanelle Pharmaceuticals Manufacturing Limited

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**Marketing authorisation date:**

28/12/2022

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**Manufacturing sites for batch release:**

Chanelle Pharmaceuticals Manufacturing Limited

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**Responsible authority:**

Federal Office Of Consumer Protection And Food Safety

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**Authorisation number:**

V7006074.00.00

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**Date of authorisation status change:**

28/12/2022

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**Reference member state:**

Ireland

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**Procedure number:**

IE/V/0770/001

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**Concerned member states:**

Austria Belgium Croatia Denmark France Germany Hungary Italy

Netherlands Poland Portugal Romania Spain

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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

English (PDF)

Published on: 8/03/2026

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Summary of Product Characteristics

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