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

Active substance:
Doramectin

Target species:
Sheep
Pig
Cattle

Route of administration:
Intramuscular use
Subcutaneous 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:
Intramuscular use:

- **Sheep**
  - Meat and offal. 70 day

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

Subcutaneous use:

- **Cattle**
  - Meat and offal. 70 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QP54AA03

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

Authorisation status:
Valid

Authorised in:
Hungary

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.

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:
Chanelle Pharmaceuticals Manufacturing Limited

Marketing authorisation date:
30/07/2025

Manufacturing sites for batch release:
Chanelle Pharmaceuticals Manufacturing Limited

Responsible authority:
Directorate Of Veterinary Medicinal Products

Authorisation number:
This information is not available for this product.

Date of authorisation status change:
30/07/2025

Reference member state:
Ireland

Procedure number:
IE/V/0770/001

Concerned member states:
Austria Belgium Croatia Denmark France Germany Hungary Italy

Netherlands Poland Portugal Romania Spain

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

Documents

Summary of Product Characteristics

Combined File of all Documents