Noromectin 10 mg/ml Solution for Injection for sheep.

Authorised

- Ivermectin

Product identification

Medicine name:
Noromectin 10 mg/ml Solution for Injection for sheep.

Active substance:
Ivermectin

Target species:
Sheep

Route of administration:
Subcutaneous use

Product details

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

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:

Subcutaneous use:

- **Sheep**
  - Meat and offal. 42 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QP54AA01

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

Authorisation status:
Valid

Authorised in:
Portugal

Available in:
Portugal

Package description:
The product will be supplied as a colourless to pale yellow solution in 1 L volume, presented in high density polyethylene vial with bromobutyl bung and aluminium cap.
The product will be supplied as a colourless to pale yellow solution in 500 ml volume, presented in high density polyethylene vial with bromobutyl bung and aluminium cap.
The product will be supplied as a colourless to pale yellow solution in 250 ml volume, presented in high density polyethylene vial with bromobutyl bung and aluminium cap.
The product will be supplied as a colourless to pale yellow solution in 100 ml volume, presented in high density polyethylene vial with bromobutyl bung and aluminium cap.
The product will be supplied as a colourless to pale yellow solution in 50 ml volume, presented in high density polyethylene vial with bromobutyl bung and aluminium cap.

Additional information

Entitlement type:
Marketing Authorisation

Marketing authorisation holder:
Norbrook Laboratories (Ireland) Limited

Marketing authorisation date:
21/09/2005

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

Responsible authority:
Directorate General For Food And Veterinary

Authorisation number:
51604

Date of authorisation status change:
10/01/2025

Reference member state:
Ireland

Procedure number:
IE/V/0168/001

Concerned member states:
Italy Portugal 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

English (PDF)
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