Noromectin 10 mg/ml Solution for Injection for Cattle and Pigs

Authorised

- Ivermectin

Product identification

Medicine name:
Noromectin 10 mg/ml Solution for Injection for Cattle and Pigs

Active substance:
Ivermectin

Target species:
Cattle
Pig

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:

- **Cattle**
  - Meat and offal. 49 day
- **Pig**
  - Meat and offal. 18 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP54AA01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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Authorisation status:

Valid

Authorised in:

Spain

Available in:

Spain

Package description:

The product will be supplied in 1 L volume, presented in high-density polyethylene vial with bromobutyl bung and aluminium cap.
The product will be supplied in 500 ml volume, presented in high-density polyethylene vial with bromobutyl bung and aluminium cap.
The product will be supplied in 250 ml volume, presented in high-density polyethylene vial with bromobutyl bung and aluminium cap.
The product will be supplied in 100 ml volume, presented in high-density polyethylene vial with bromobutyl bung and aluminium cap.
The product will be supplied in 50 ml volume, presented in high-density polyethylene vial with bromobutyl bung and aluminium cap.

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:
Norbrook Laboratories (Ireland) Limited

Marketing authorisation date:
13/11/2000

Manufacturing sites for batch release:
Norbrook Laboratories Limited
Norbrook Laboratories (Ireland) Limited

Responsible authority:
Spanish Agency Of Medicines And Medical Devices

Authorisation number:
1352 ESP

Date of authorisation status change:
26/02/2018

Reference member state:
Ireland

Procedure number:
IE/V/0104/001

Concerned member states:
France Germany Greece Iceland Italy Netherlands Portugal Spain

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

Documents

Summary of Product Characteristics

English (PDF)
Published on: 29/12/2024
Download

Labelling

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.

Combined File of all Documents

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