

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

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Pig

- Meat and offal. 18 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP54AA01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Netherlands

Available in:

Netherlands

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:

14/08/2003

Manufacturing sites for batch release:

Norbrook Laboratories Limited

Norbrook Laboratories (Ireland) Limited

Responsible authority:

Medicines Evaluation Board

Authorisation number:

REG NL 10107

Date of authorisation status change:

25/01/2022

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

Combined File of all Documents

English (PDF)

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Summary of Product Characteristics

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