

Noromectin Multi Injection Solution for Injection

Authorised

- Ivermectin

Product identification

Medicine name:

Noromectin Multi Injection Solution for Injection

Active substance:

Ivermectin

Target species:

Cattle

Sheep

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:

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Cattle

- Meat and offal. 49 day

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Sheep

- Meat and offal. 42 day

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Pig

- Meat and offal. 28 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP54AA01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Ireland

Available in:

Ireland

Package description:

Noromectin Multi Injection will be supplied in a 1.5 litre (1 x 1 litre and 1 x 500ml) combination pack.

Noromectin Multi Injection will be supplied in 50 ml volume, presented in high density polyethylene vial with bromobutyl bung and aluminium cap.

Noromectin Multi Injection will be supplied in 100 ml volume, presented in high density polyethylene vial with bromobutyl bung and aluminium cap.

Noromectin Multi Injection will be supplied in 250 ml volume, presented in high density polyethylene vial with bromobutyl bung and aluminium cap.

Noromectin Multi Injection will be supplied in 500 ml volume, presented in high density polyethylene vial with bromobutyl bung and aluminium cap.

Noromectin Multi Injection will be supplied in 1 litre volume, presented in high density polyethylene vial with bromobutyl bung and aluminium cap.

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Informed consent (abridged application) - Council Directive 81/851/EEC

Marketing authorisation holder:

Norbrook Laboratories (Ireland) Limited

Marketing authorisation date:

22/02/2002

Manufacturing sites for batch release:

Norbrook Manufacturing Limited

Norbrook Laboratories Limited

Responsible authority:

Health Products Regulatory Authority

Authorisation number:

VPA22664/068/001

Date of authorisation status change:

22/02/2002

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