

Noromectin vet , 10 mg/ml, solution for injection

Authorised

- Ivermectin

Product identification

Medicine name:

Noromectin vet , 10 mg/ml, solution for injection

Active substance:

Ivermectin

Target species:

Cattle

Reindeer

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

Lactating dairy cattle must not be injected. Dry cows and heifers must not be injected later than 60 days prior to calving.

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Reindeer

- Meat and offal. 28 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:

Sweden

Available in:

Sweden

Package description:

Available only in Swedish

Available only in Swedish

Available only in Swedish

Available only in Swedish

Available only in Swedish

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Well-established use application (Article 13a of Directive No 2001/82/EC)

Marketing authorisation holder:

Norbrook Laboratories (Ireland) Limited

Marketing authorisation date:

23/02/2001

Manufacturing sites for batch release:

Norbrook Manufacturing Limited

Norbrook Laboratories Limited

Responsible authority:

Swedish Medical Products Agency

Authorisation number:

14895

Date of authorisation status change:

23/02/2001

Reference member state:

Sweden

Procedure number:

SE/V/0108/001

Concerned member states:

Finland

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

Documents

Package Leaflet

English (PDF)

Published on: 26/04/2022

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Summary of Product Characteristics

English (PDF)

Published on: 26/04/2022

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