

Noromectin pour-on vet, 5 mg/ml, pour-on solution

Authorised

- Ivermectin

Product identification

Medicine name:

Noromectin pour-on vet, 5 mg/ml, pour-on solution

Active substance:

Ivermectin

Target species:

Cattle

Route of administration:

Pour-on use

Product details

Active substance and strength:

Ivermectin

5.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Pour-on solution

Withdrawal period by route of administration:**Pour-on use:**

-

Cattle

- Meat and offal. 21 day

Lactating dairy cattle must not be treated. Dry cows or heifers must not be treated later than 60 days before calving.

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP54AA01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Finland

Available in:

Finland

Package description:

Bottle, 1000 ml (collapsible)

Bottle, 1000 ml (collapsible)

Bottle, 1000 ml (collapsible)

Bottle, 1000 ml (collapsible)

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:

Norbrook Laboratories (Ireland) Limited

Marketing authorisation date:

27/02/2002

Manufacturing sites for batch release:

Norbrook Manufacturing Limited

Norbrook Laboratories Limited

Responsible authority:

Finnish Medicines Agency

Authorisation number:

13841

Date of authorisation status change:

27/02/2002

Reference member state:

Sweden

Procedure number:

SE/V/0107/001

Concerned member states:

Finland

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

Documents

Summary of Product Characteristics

English (PDF)

Published on: 26/02/2026

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Package Leaflet

English (PDF)

Published on: 26/02/2026

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Labelling