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Macromectin 0.8 mg/ml oral solution for sheep

Authorised

- Ivermectin

Product identification

Medicine name:
Macromectin 0.8 mg/ml oral solution for sheep

Active substance:
Ivermectin

Target species:
Sheep

Route of administration:
Oral use

Product details

Active substance and strength:
Ivermectin
0.80 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Oral solution

Withdrawal period by route of administration:

Oral use:

- **Sheep**

  - Meat and offal. 10 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:

Sweden

Available in:

Sweden

Package description:

The product will be supplied in 1.0 L high density polyethylene jerry-can containers complete with polypropylene caps

The product will be supplied in 5.0 L high density polyethylene jerry-can containers complete with polypropylene caps

The product will be supplied in 2 x 5.0 L high density polyethylene jerry-can containers complete with polypropylene caps

The product will be supplied in 2.5 L high density polyethylene back-pack containers complete with polypropylene plastic screw caps.

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The product will be supplied in 2.5 L high density polyethylene jerry-can containers complete with polypropylene caps

The product will be supplied in 1.0 L high density polyethylene back-pack containers complete with polypropylene plastic screw caps.

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:
8/12/2006

Manufacturing sites for batch release:
Norbrook Manufacturing Limited
Norbrook Laboratories Limited

Responsible authority:
Swedish Medical Products Agency

Authorisation number:
23313

Date of authorisation status change:
8/12/2006

Reference member state:
Ireland

Procedure number:
IE/V/0179/001

Concerned member states:
Sweden

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

Documents

Summary of Product Characteristics

English (PDF)
Published on: 18/01/2026
Updated on: 14/03/2026
Download

Package Leaflet

This document does not exist in this language (English). You can find it in another language below.

Combined File of all Documents

Labelling

This document does not exist in this language (English). You can find it in another language below.