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
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Package Leaflet

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

Combined File of all Documents