Macromectin 0.8 mg/ml oral solution for sheep

Authorised

- Ivermectin

Product identification

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

Package description:
The product will be supplied in 1.0 L high density polyethylene back-pack containers complete with polypropylene plastic screw caps.
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 5.0 L high density polyethylene back-pack containers complete with polypropylene plastic screw caps.
The product will be supplied in 2 x 5.0 L high density polyethylene back-pack containers complete with polypropylene plastic screw caps.
The product will be supplied in 2.5 L high density polyethylene back-pack containers complete with polypropylene plastic screw 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 5.0 L high density polyethylene jerry-can containers complete with polypropylene caps
The product will be supplied in 1.0 L high density polyethylene jerry-can containers complete with polypropylene caps

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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:
2/09/2005

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

Responsible authority:
Health Products Regulatory Authority

Authorisation number:
VPA22664/075/001

Date of authorisation status change:
2/09/2005

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
