

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

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

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 Manufacturing Limited

Norbrook Laboratories 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

Combined File of all Documents