Ecomectin 5 mg/ml Pour-on Solution for Cattle

Authorised

- Ivermectin

Product identification

Medicine name:
Ecomectin 5 mg/ml Pour-on Solution for Cattle
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Active substance:
Ivermectin

Target species:
Cattle

Route of administration:
Topical 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:

Topical use:

- **Cattle**

  - Meat and offal. 31 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QP54AA01

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
United Kingdom (Northern Ireland)

Package description:
5.0 L white non-fluorinated or fluorinated high-density polyethylene back-pack with polypropylene strap and vented cap Closure: White polypropylene screw-cap.
2.5 L white non-fluorinated or fluorinated high-density polyethylene back-pack with polypropylene strap and vented cap Closure: White polypropylene screw-cap.
1.0 L white non-fluorinated or fluorinated high-density polyethylene bottle with a drawing tube and measuring device. Closure: White polypropylene screw-cap.
250 ml white non-fluorinated or fluorinated high-density polyethylene bottle with a drawing tube and measuring device. Closure: White polypropylene screw-cap.

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:
Eco Animal Health Limited

Marketing authorisation date:

6/08/2007

Manufacturing sites for batch release:
Acme Drugs - S.r.l.
Safapac (Cambridge) Limited

Responsible authority:
The Veterinary Medicines Directorate

Authorisation number:
Vm 13277/4021

Date of authorisation status change:
22/08/2022

Reference member state:
Ireland

Procedure number:
IE/V/0108/001

Concerned member states:
Belgium France Germany Greece Italy Portugal
United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics
