Macromectin 5 mg/ml Pour-On Solution

Authorised

- Ivermectin

Product identification

Medicine name:
Macromectin 5 mg/ml Pour-On Solution
BAYMEC 5 MG/ML SOLUTION POUR-ON

Active substance:
Ivermectin

Target species:
Cattle

Route of administration:
Pour-on 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:

Pour-on use:

- **Cattle**
  - Meat and offal. 28 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:

France

Available in:

France

Package description:

The product will be supplied in: 250 ml twin-neck high density polyethylene dispensers
The product will be supplied in:1.0 L single-neck high density polyethylene dispensers
The product will be supplied in:2.5 L low density polyethylene backpacks.
The product will be supplied in:5 L low density polyethylene backpacks.
The product will be supplied in: 250 ml squeeze-measure high density polyethylene dispensers
The product will be supplied in:1.0 L squeeze-measure high density polyethylene dispensers
The product will be supplied in:250 ml single-neck high density polyethylene dispensers
The product will be supplied in:1 L high density polyethylene backpacks.
The product will be supplied in: 1.0 L twin-neck high density polyethylene dispensers

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Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:
Norbrook Laboratories (Ireland) Limited

Marketing authorisation date:
11/05/2006

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

Responsible authority:
French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:
FR/V/0681919 0/2006

Date of authorisation status change:
11/05/2011

Reference member state:
Ireland

Procedure number:
IE/V/0180/001

Concerned member states:
France

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

Documents

Summary of Product Characteristics

English (PDF)
Published on: 6/07/2025
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Combined File of all Documents

Package Leaflet and Labelling

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