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Ridamec 1 mg/ml oral solution for sheep

Authorised

- Moxidectin

Product identification

Medicine name:
Ridamec 1 mg/ml oral solution for sheep

Active substance:
Moxidectin

Target species:
Sheep

Route of administration:
Oral use

Product details

Active substance and strength:
Moxidectin
1.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Oral solution

Withdrawal period by route of administration:

Oral use:

- **Sheep**
  - Meat and offal. 14 day
  - Milk. 5 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP54AB02

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Norway

Package description:

White HDPE flexi containers containing 5 L of product. Thecontainers are closed with an aluminium foil seal and polypropylene tamper-evidentcaps. The product is marketed in a cardboard outer carton.
White HDPE flexi containers containing 3 L of product. Thecontainers are closed with an aluminium foil seal and polypropylene tamper-evidentcaps. The product is marketed in a cardboard outer carton.
White HDPE flexi containers containing 2.5 L of product. Thecontainers are closed with an aluminium foil seal and polypropylene tamper-evidentcaps. The product is marketed in a cardboard outer carton.
White HDPE flexi containers containing 1L of product. Thecontainers are closed with an aluminium foil seal and polypropylene tamper-evidentcaps. The product is marketed in a cardboard outer carton.

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:
Chanelle Pharmaceuticals Manufacturing Limited

Marketing authorisation date:
25/04/2019

Manufacturing sites for batch release:
Chanelle Pharmaceuticals Manufacturing Limited

Responsible authority:
Norwegian Medical Products Agency

Authorisation number:
17-11982

Date of authorisation status change:
25/04/2019

Reference member state:
Ireland

Procedure number:
IE/V/0403/001

Concerned member states:
Austria Belgium Bulgaria Croatia Cyprus Denmark France Greece Hungary Iceland Italy Luxembourg Netherlands Norway Poland Portugal Sweden United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics

English (PDF)
Published on: 18/05/2025
Updated on: 13/03/2026
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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