Tilmodil 300 mg/ml Solution for Injection for cattle and sheep

Authorised

- Tilmicosin

Product identification

Medicine name:
Tilmodil 300 mg/ml Solution for Injection for cattle and sheep

Active substance:
Tilmicosin

Target species:
Cattle
Sheep

Route of administration:
Subcutaneous use

Product details

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

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:

Subcutaneous use:

- **Cattle**
  - Meat and offal. 70 day
  - Milk. 36 day
- **Sheep**
  - Meat and offal. 42 day
  - Milk. 18 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01FA91

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

---

Authorisation status:

Valid

Authorised in:

Netherlands

Package description:

Cardboard box with 1 multi-dose amber coloured, glass vial (type II) of 50 ml with bromobutyl, tefloncoated stopper and an aluminium overseal.
Cardboard box with 12 multi-dose amber coloured, glass vials (type II) of 50 ml with bromobutyl, tefloncoated stoppers and an aluminium overseal.
Cardboard box with 1 multi-dose amber coloured, glass vial (type II) of 100 ml with bromobutyl, tefloncoated stopper and an aluminium overseal.
Cardboard box with 12 multi-dose amber coloured, glass vials (type II) of 100 ml with bromobutyl, tefloncoated stoppers and an aluminium overseal.

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:
Emdoka

Marketing authorisation date:
10/03/2011

Manufacturing sites for batch release:
Produlab Pharma B.V.

Responsible authority:
Medicines Evaluation Board

Authorisation number:
REG NL 106754

Date of authorisation status change:
13/06/2024

Reference member state:
Ireland

Procedure number:
IE/V/0564/001

Concerned member states:
Belgium Germany Luxembourg Netherlands United Kingdom (Northern Ireland)

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

Documents

Combined File of all Documents

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

Summary of Product Characteristics