HYMATIL 300 mg/ml Solution for injection for cattle

Authorised

- Tilmicosin

Product identification

Medicine name:
HYMATIL 300 mg/ml Solution for injection for cattle
Hymatil 300 mg/ml Roztwór do wstrzykiwań

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

- **Sheep**
  - Meat and offal. 42 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01FA91

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Poland

Package description:

box containing 12 vials of 250 ml
box containing 12 vials of 100 ml
box containing 12 vials of 50 ml
box containing 10 vials of 250 ml
box containing 10 vials of 100 ml
box containing 10 vials of 50 ml
box containing 6 vials of 250 ml
box containing 6 vials of 100 ml
box containing 6 vials of 50 ml
box containing 1 vial of 250 ml
box containing 1 vial of 100 ml
box containing 1 vial of 50 ml

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:
Industrial Veterinaria S.A.

Marketing authorisation date:
9/06/2010

Manufacturing sites for batch release:
Industrial Veterinaria S.A.
aniMedica GmbH

Responsible authority:
Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

Authorisation number:
1987

Date of authorisation status change:
9/06/2010

Reference member state:
Spain

Procedure number:
ES/V/0141/001

Concerned member states:
Austria Belgium Bulgaria Cyprus Czechia Estonia Germany Greece Hungary Ireland Italy Latvia Lithuania Poland Portugal Romania Slovakia Slovenia United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics

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

Labelling

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

Package Leaflet

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