

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 šķīdums injekcijām liellopiem un aitām

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:

Latvia

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:

21/09/2009

Manufacturing sites for batch release:

Industrial Veterinaria S.A.

Animedica GmbH

Responsible authority:

Food And Veterinary Service

Authorisation number:

V/DCP/09/0021

Date of authorisation status change:

21/09/2009

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

Labelling

Source URL: <https://medicines.health.europa.eu/veterinary/600000015587>