Tullavis 100 mg/ml solution for injection for cattle, pigs and sheep

Authorised

- Tulathromycin

Product identification

Medicine name:
Tullavis 100 mg/ml solution for injection for cattle, pigs and sheep

Active substance:
Tulathromycin

Target species:
Sheep
Pig
Cattle

Route of administration:
Intramuscular use
Subcutaneous use

Product details

Active substance and strength:
Tulathromycin

100.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:
Intramuscular use:

- **Sheep**
  - Meat and offal. 16 day
- **Pig**
  - Meat and offal. 13 day

Subcutaneous use:

- **Cattle**
  - Meat and offal. 22 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01FA94

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Germany

Available in:
Germany

Package description:
Clear glass (Type II Ph. Eur.) vial closed with a bromobutyl rubber stopper and sealed with an aluminium cap.Cardboard box containing 1 vial of 20 ml.

Clear glass (Type II Ph. Eur.) vial closed with a bromobutyl rubber stopper and sealed with an aluminium cap.Cardboard box containing 1 vial of 50 ml.
Clear glass (Type II Ph. Eur.) vial closed with a bromobutyl rubber stopper and sealed with an aluminium cap.Cardboard box containing 1 vial of 100 ml.
Clear glass (Type II Ph. Eur.) vial closed with a bromobutyl rubber stopper and sealed with an aluminium cap.Cardboard box containing 1 vial of 250 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:
27/08/2020

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

Responsible authority:
Federal Office Of Consumer Protection And Food Safety

Authorisation number:
402709.00.00

Date of authorisation status change:
27/08/2020

Reference member state:
Ireland

Procedure number:

IE/V/0893/001

Concerned member states:

Austria Belgium Bulgaria Croatia Cyprus Czechia Denmark Estonia Finland France Germany Greece Hungary Iceland Italy Latvia Lithuania Netherlands Poland Portugal Romania Slovakia Slovenia Spain

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: 21/12/2023
Download

Combined File of all Documents

English (PDF)
Published on: 9/11/2025
Download

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