Zactran 150 mg/ml - Solution for injection (cattle, pigs, sheep)

Authorised

- Gamithromycin

Product identification

Medicine name:
Zactran 150 mg/ml - Solution for injection (cattle, pigs, sheep)

Active substance:
Gamithromycin

Target species:
Cattle
Sheep
Pig

Route of administration:
Intramuscular use
Subcutaneous use

Product details

Active substance and strength:
Gamithromycin
150.00 milligram(s) / 1.00 Vial

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Cattle**
  - Meat and offal. 64 day 64 days
- **Sheep**
  - Meat and offal. 29 day 29 days
- **Pig**
  - Meat and offal. 16 day 16 days

Subcutaneous use:

- **Cattle**
  - Meat and offal. 64 day 64 days
- **Sheep**
  - Meat and offal. 29 day 29 days
- **Pig**
  - Meat and offal. 16 day 16 days

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01FA95

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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Authorisation status:

Valid

Authorised in:

Austria , Belgium , Bulgaria , Croatia , Cyprus , Czechia , Denmark , Estonia , Finland , France , Germany , Greece , Hungary , Iceland , Ireland , Italy , Latvia , Liechtenstein , Lithuania , Luxembourg , Malta , Netherlands , Norway , Poland , Portugal , Romania , Slovakia , Slovenia , Spain , Sweden , United Kingdom (Northern Ireland)

Package description:

Packaging:Vial (glass), Package_size:1 vial, Content:50 ml

Packaging:Vial (PP), Package_size:1 vial, Content:250 ml

Packaging:Vial (PP), Package_size:1 vial, Content:100 ml
Packaging:Vial (glass), Package_size:1 vial, Content:250 ml
Packaging:Vial (glass), Package_size:1 vial, Content:100 ml

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Boehringer Ingelheim Vetmedica GmbH

Marketing authorisation date:
24/07/2008

Manufacturing sites for batch release:
Boehringer Ingelheim Animal Health France

Responsible authority:
European Commission

Authorisation number:
This information is not available for this product.

Date of authorisation status change:
24/07/2008

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

Documents

Combined File of all Documents

English (PDF)
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