Draxxin 100 mg/ml - Solution for injection (cattle, pig, sheep)

Authorised

- Tulathromycin

Product identification

Medicine name:
Draxxin 100 mg/ml - Solution for injection (cattle, pig, sheep)

Active substance:
Tulathromycin

Target species:
Cattle
Sheep
Pig

Route of administration:
Intramuscular use
Subcutaneous use

Product details

Active substance and strength:
Tulathromycin
100.00 milligram(s) / 1.00 Vial

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Cattle**
  - Meat and offal. 22 day — 22 days
- **Sheep**
  - Meat and offal. 16 day — 16 days
- **Pig**
  - Meat and offal. 13 day — 13 days

Subcutaneous use:

- **Cattle**
  - Meat and offal. 22 day — 22 days
- **Sheep**
  - Meat and offal. 16 day — 16 days
- **Pig**
  - Meat and offal. 13 day — 13 days

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01FA94

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)

Available in:
Belgium , Bulgaria , Croatia , Cyprus , Czechia , Denmark , Estonia , Finland , Greece , Hungary , Ireland , Italy , Latvia , Lithuania , Luxembourg , Netherlands , Poland , Romania , Slovakia , Spain , Sweden , United Kingdom (Northern Ireland)

Package description:
Packaging:Vial (glass), Package_size:1 vial, Content:250 ml
Packaging:Vial (glass), Package_size:1 vial, Content:100 ml
Packaging:Vial (glass), Package_size:1 vial, Content:50 ml
Packaging:Vial (glass), Package_size:1 vial, Content:20 ml

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Complete application (stand-alone) - Council Directive 81/851/EEC

Marketing authorisation holder:
Zoetis Belgium SA

Marketing authorisation date:
11/11/2003

Manufacturing sites for batch release:
Fareva Amboise
Zoetis Belgium
Zoetis Manufacturing & Research Spain S.L.

Responsible authority:
European Commission

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

Date of authorisation status change:
11/11/2003

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

Documents

Combined File of all Documents

English (PDF)
Published on: 18/04/2024
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