

Tulissin 100 mg/ml - Solution for injection (cattle, pigs, sheep)

Authorised

- Tulathromycin

Product identification

Medicine name:

Tulissin 100 mg/ml - Solution for injection (cattle, pigs, 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

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:

Austria , Belgium , Croatia , Czechia , Denmark , Estonia , Finland , France , Germany , Greece , Hungary , Ireland , Italy , Latvia , Lithuania , Poland , Portugal , Romania , Slovakia , Slovenia , Spain , United Kingdom (Northern Ireland)

Package description:

Packaging:Vial (glass), Package_size:1 vial, Content:20 ml

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

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

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

Packaging:Vial (glass) with protective sleeve, Package_size:1 vial, Content: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:

Virbac

Marketing authorisation date:

24/04/2020

Manufacturing sites for batch release:

Fareva Amboise

Virbac

Responsible authority:

European Commission

Authorisation number:

This information is not available for this product.

Date of authorisation status change:

24/04/2020

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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