

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

Authorised

- Tulathromycin

Product identification

Medicine name:

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

Active substance:

Tulathromycin

Target species:

Cattle

Route of administration:

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:

Subcutaneous use:

-

Cattle

- Meat and offal. 22 day 22 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)

Package description:

Packaging:Vial (glass), Package_size:1 vial, Content:500 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:

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)

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