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DRALTEN 100 MG/ML SOLUTION INJECTABLE POUR BOVINS, PORCINS ET OVINS

Not
authorised

- Tulathromycin

Product identification

Medicine name:

DRALTEN 100 MG/ML SOLUTION INJECTABLE POUR BOVINS, PORCINS ET OVINS

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:

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Sheep

- Milk. no withdrawal period No withdrawal period
- Meat and offal. 16 day

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Pig

- Meat and offal. 13 day

Subcutaneous use:

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Cattle

- Milk. no withdrawal period No withdrawal period
- Meat and offal. 22 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01FA94

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription except for some pack sizes

Authorisation status:

Surrendered

Authorised in:

France

Package description:

Available only in French

Available only in [French](#)

Available only in [French](#)

Available only in [French](#)

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:

Vetoquinol S.A.

Marketing authorisation date:

25/09/2020

Manufacturing sites for batch release:

Vetoquinol S.A.

Responsible authority:

French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:

FR/V/5259747 1/2020

Date of authorisation status change:

26/08/2025

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

Documents

Summary of Product Characteristics

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