

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

Authorised

- Gamithromycin

## Product identification

**Medicine name:**

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

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**Active substance:**

Gamithromycin

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**Target species:**

Cattle

Pig

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**Route of administration:**

Intramuscular use

Subcutaneous use

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## Product details

**Active substance and strength:**

Gamithromycin

150.00 milligram(s) / 1.00 Vial

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**Pharmaceutical form:**

Solution for injection

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**Withdrawal period by route of administration:**

**Intramuscular use:**

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**Cattle**

- Meat and offal. 64 day 64 days

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**Pig**

- Meat and offal. 16 day 16 days

**Subcutaneous use:**

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**Cattle**

- Meat and offal. 64 day 64 days

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**Pig**

- Meat and offal. 16 day 16 days

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QJ01FA95

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**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)

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**Package description:**

Packaging:Vial (PP), Package\_size:1 vial, Content:500 ml  
Packaging:Vial (glass), Package\_size:1 vial, Content:500 ml

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Full application (Article 12(3) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Boehringer Ingelheim Vetmedica GmbH

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**Marketing authorisation date:**

24/07/2008

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**Manufacturing sites for batch release:**

Fresenius Kabi Austria GmbH

Boehringer Ingelheim Animal Health France SCS

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**Responsible authority:**

European Commission

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**Authorisation number:**

This information is not available for this product.

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**Date of authorisation status change:**

20/07/2009

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To consult adverse reactions on veterinary medicinal products please go to  
[www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Combined File of all Documents

English (PDF)

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