

# COBACTAN LA 7.5% w/v suspension for injection for cattle

Authorised

- Cefquinome sulfate

## Product identification

**Medicine name:**

COBACTAN LA 7.5% w/v suspension for injection for cattle

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**Active substance:**

Cefquinome sulfate

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**Target species:**

Cattle

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**Route of administration:**

Subcutaneous use

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## Product details

**Active substance and strength:**

Cefquinome sulfate

88.90 milligram(s) / 1.00 millilitre(s)

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**Pharmaceutical form:**

Suspension for injection

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**Withdrawal period by route of administration:****Subcutaneous use:**

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**Cattle**

- Meat and offal. 13 day
- Milk. no withdrawal period

Do not use in dairy cows producing milk for human consumption (during lactation or the dry period). Do not use within two months prior to first calving in heifers intended for the production of milk for human consumption.

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

QJ01DE90

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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:**

Germany

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**Available in:**

Germany

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**Package description:**

(ID1) 100 millilitre(s): unspecified outer container with 1 Vial (glass) with 100 millilitre(s)

(ID2) 250 millilitre(s): unspecified outer container with 1 Vial (glass) with 250 millilitre(s)

(ID3) 50 millilitre(s): unspecified outer container with 1 Vial (glass) with 50 millilitre(s)

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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:**

Intervet Deutschland GmbH

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**Marketing authorisation date:**

28/07/2006

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**Manufacturing sites for batch release:**

Intervet International GmbH

Intervet Productions S.r.l.

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**Responsible authority:**

Federal Office Of Consumer Protection And Food Safety

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**Authorisation number:**

400944.00.00

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**Date of authorisation status change:**

20/06/2011

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**Reference member state:**

Germany

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**Procedure number:**

DE/V/0145/001

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**Concerned member states:**

Austria Belgium Cyprus Czechia Denmark France Greece Hungary Italy  
Luxembourg Netherlands Poland Portugal Slovakia Slovenia Spain

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## Documents

Combined File of all Documents

English (PDF)

Published on: 21/08/2025

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