

# Ceftiomax 50 mg/ml suspension for injection for swine and cattle

Authorised

- Ceftiofur

## Product identification

**Medicine name:**

Ceftiomax 50 mg/ml suspension for injection for swine and cattle

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**Active substance:**

Ceftiofur

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**Target species:**

Pig

Cattle

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**Route of administration:**

Intramuscular use

Subcutaneous use

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## Product details

**Active substance and strength:**

Ceftiofur

50.00 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:**

**Intramuscular use:**

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**Pig**

- Meat and offal. 5 day

**Subcutaneous use:**

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**Cattle**

- Milk. 0 day

- Meat and offal. 8 day

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

QJ01DD90

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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:**

Poland

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**Package description:**

Clear glass vial type I of 250 ml with a bromobuthyl pink rubber stopper and aluminium cap gold colour. One vial of 250ml is available in a cardboard box.

Clear glass vial type I of 100 ml with a bromobuthyl rubber stopper and aluminium cap with opening ring FLIPP OFF of blue colour. One vial of 100ml is available in a cardboard box.

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Generic application (Article 13(1) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Laboratorios Calier S.A.

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**Marketing authorisation date:**

20/10/2008

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**Manufacturing sites for batch release:**

Laboratorios Calier S.A.

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**Responsible authority:**

Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

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**Authorisation number:**

1859

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**Date of authorisation status change:**

20/10/2008

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**Reference member state:**

Portugal

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**Procedure number:**

PT/V/0101/001

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**Concerned member states:**

Austria Belgium Bulgaria Czechia Estonia France Germany Greece Hungary  
Italy Latvia Lithuania Netherlands Poland Romania Slovakia Spain  
United Kingdom (Northern Ireland)

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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

### Summary of Product Characteristics

English (PDF)

Published on: 28/05/2026

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### Labelling

This document does not exist in this language (English). You can find it in another language below.

### Package Leaflet

This document does not exist in this language (English). You can find it in another language below.