

# 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

---

**Active substance:**

Ceftiofur

---

**Target species:**

Pig

Cattle

---

**Route of administration:**

Intramuscular use

Subcutaneous use

---

## Product details

**Active substance and strength:**

Ceftiofur

50.00 milligram(s) / 1.00 millilitre(s)

---

**Pharmaceutical form:**

Suspension for injection

---

**Withdrawal period by route of administration:**

**Intramuscular use:**

- 

**Pig**

- Meat and offal. 5 day

**Subcutaneous use:**

- 

**Cattle**

- Milk. 0 day

- Meat and offal. 8 day

---

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

QJ01DD90

---

**Legal status of supply:**

Veterinary medicinal product not subject to veterinary prescription

---

**Authorisation status:**

Valid

---

**Authorised in:**

Estonia

---

**Package description:**

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.

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.

---

## 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:**

Laboratorios Calier S.A.

---

**Marketing authorisation date:**

23/05/2010

---

**Manufacturing sites for batch release:**

Laboratorios Calier S.A.

---

**Responsible authority:**

State Agency Of Medicines

---

**Authorisation number:**

1602

---

**Date of authorisation status change:**

23/05/2010

---

**Reference member state:**

Portugal

---

**Procedure number:**

PT/V/0101/001

---

**Concerned member states:**

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

---

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: 21/08/2024

[Download](#)