# CEFOKEL 50 mg/ml, suspension for injection for pigs and cattle

Authorised

- Ceftiofur hydrochloride

## Product identification

**Medicine name:**
CEFOKEL 50 mg/ml, suspension for injection for pigs and cattle
CEFOKEL

**Active substance:**
Ceftiofur hydrochloride

**Target species:**
Pig
Cattle

**Route of administration:**
Intramuscular use
Subcutaneous use

## Product details

**Active substance and strength:**
Ceftiofur hydrochloride
53.48 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**
  - Meat and offal. 8 day
  - Milk. 0 hour

**Anatomical therapeutic chemical veterinary (ATCvet) codes:**
QJ01DD90

**Legal status of supply:**
Veterinary medicinal product subject to veterinary prescription

**Authorisation status:**
Valid

**Authorised in:**
Germany

**Available in:**
Germany

**Package description:**
Colourless glass type I vial of 100 ml, closed with grey coated bromobutyl rubber stoppers and aluminium caps.Vials are individually packed in a carton box.Twelve vials are grouped as a clinical pack.
Colourless glass type I vial of 100 ml, closed with grey coated bromobutyl rubber stoppers and aluminium caps.Vials are individually packed in a carton box.Ten vials are grouped as a clinical pack.

Colourless glass type I vial of 100 ml, closed with grey coated bromobutyl rubber stoppers and aluminium caps.Vials are individually packed in a carton box.Six vials are grouped as a clinical pack.
Colourless glass type I vial of 100 ml, closed with grey coated bromobutyl rubber stopper and aluminium cap.Vial is individually packed in a carton box.One vial is grouped as a clinical pack.

## 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:**
KELA Kempisch Laboratorium Kela Laboratoria

**Marketing authorisation date:**
16/04/2013

**Manufacturing sites for batch release:**
KELA Kempisch Laboratorium Kela Laboratoria

**Responsible authority:**
Federal Office Of Consumer Protection And Food Safety

**Authorisation number:**
401801.00.00

**Date of authorisation status change:**
31/10/2018

**Reference member state:**
Ireland

**Procedure number:**
IE/V/0303/001

**Concerned member states:**
Belgium Germany Luxembourg Poland Romania Spain
United Kingdom (Northern Ireland)

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

## Documents

Summary of Product Characteristics

English (PDF)
Published on: 28/09/2025
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