

# DIACEF 50 mg/ml

Authorised

- Ceftiofur hydrochloride

## Product identification

**Medicine name:**

DIACEF 50 mg/ml

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**Active substance:**

Ceftiofur hydrochloride

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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 hydrochloride

53.48 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**

- Meat and offal. 6 day

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**Cattle**

- Milk. 0 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:**

Romania

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**Available in:**

Romania

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**Package description:**

cardboard box containing 1 vial of 250 ml

cardboard box containing 1 vial of 100 ml

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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:**

Super's Diana S.L.

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**Marketing authorisation date:**

10/06/2013

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**Manufacturing sites for batch release:**

Super's Diana S.L.

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**Responsible authority:**

Institute For Control Of Biological Products And Veterinary Medicines

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**Authorisation number:**

180089

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**Date of authorisation status change:**

3/11/2025

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**Reference member state:**

Spain

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**Procedure number:**

ES/V/0187/001

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**Concerned member states:**

Czechia Hungary Poland Romania

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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

Package Leaflet

Combined File of all Documents

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