

Ceftionil 50 mg/ml suspension for injection for pigs and cattle

Authorised

- Ceftiofur hydrochloride

Product identification

Medicine name:

Ceftionil 50 mg/ml suspension for injection for pigs and cattle

Active substance:

Ceftiofur hydrochloride

Target species:

Cattle

Pig

Route of administration:

Subcutaneous use

Intramuscular 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:

Subcutaneous use:

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Cattle

- Meat and offal. 8 day
- Milk. 0 hour

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Cattle

- Meat and offal. 8 day
- Milk. 0 hour

Intramuscular use:

-

Pig

- Meat and offal. 5 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01DD90

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Portugal

Package description:

box containing 1 vial of 250 ml

box containing 1 vial of 100 ml

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 E Industrias Iven S.A.

Marketing authorisation date:

19/07/2011

Manufacturing sites for batch release:

Laboratorios Maymo S.A.U.

Responsible authority:

Directorate General For Food And Veterinary

Authorisation number:

358/01/11DFVPT

Date of authorisation status change:

13/07/2026

Reference member state:

Spain

Procedure number:

ES/V/0165/001

Concerned member states:

Czechia Hungary Italy Poland Portugal Slovakia

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