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

Authorised

- Ceftiofur hydrochloride

Product identification

Medicine name:
ACTIONIS 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
50.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Suspension for injection

Withdrawal period by route of administration:

Subcutaneous use:

- **Cattle**
  - Meat and offal. 6 day
  - Milk. 0 hour
  - Meat and offal. 6 day
  - Milk. 0 hour
- **Cattle**
  - Meat and offal. 6 day
  - Milk. 0 hour
  - Meat and offal. 6 day
  - Milk. 0 hour

Intramuscular use:

- **Pig**
  - Meat and offal. 6 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

Available in:

Portugal

Package description:
box containing 1 PET vial of 250 ml
box containing 1 PET 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 Syva S.A.

Marketing authorisation date:
3/03/2011

Manufacturing sites for batch release:
Laboratorios Syva S.A.

Responsible authority:
Directorate General For Food And Veterinary

Authorisation number:
324/01/11DFVPT

Date of authorisation status change:
11/10/2023

Reference member state:
Spain

Procedure number:
ES/V/0157/001

Concerned member states:

Germany Hungary Italy Poland Portugal United Kingdom (Northern Ireland)

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

Documents

Combined File of all Documents

English (PDF)
Published on: 24/07/2025
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Summary of Product Characteristics

Package Leaflet

Labelling