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:

United Kingdom (Northern Ireland)

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:
7/06/2011

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

Responsible authority:
The Veterinary Medicines Directorate

Authorisation number:
Vm 31592/3003

Date of authorisation status change:
4/05/2024

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

Summary of Product Characteristics

Package Leaflet

Labelling

Combined File of all Documents