Ceftiomax 50 mg/ml suspension for injection for swine and cattle

Authorised

- Ceftiofur

Product identification

Medicine name:
Ceftiomax 50 mg/ml suspension for injection for swine and cattle
Ceftiomax 50 mg/ml injekcinė suspensija kiaulėms ir galvijams

Active substance:
Ceftiofur

Target species:
Pig
Cattle

Route of administration:
Intramuscular use
Subcutaneous use

Product details

Active substance and strength:
Ceftiofur
50.00 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**
  - Milk. 0 day
  - Meat and offal. 8 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01DD90

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

Authorisation status:
Valid

Authorised in:
Lithuania

Package description:
Clear glass vial type I of 100 ml with a bromobuthyl rubber stopper and aluminium cap with opening ring FLIPP OFF of blue colour. One vial of 100ml is available in a cardboard box.
Clear glass vial type I of 250 ml with a bromobuthyl pink rubber stopper and aluminium cap gold colour. One vial of 250ml is available in a cardboard box.

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 Calier S.A.

Marketing authorisation date:
10/04/2010

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

Responsible authority:
State Food And Veterinary Service

Authorisation number:
LT/2/10/1928/001-002

Date of authorisation status change:
17/12/2013

Reference member state:
Portugal

Procedure number:
PT/V/0101/001

Concerned member states:
Austria Belgium Bulgaria Czechia Estonia France Germany Greece Hungary Italy Latvia Lithuania Netherlands Poland Romania Slovakia 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

RV1928.pdf