CEFTIONIL

Authorised

- Ceftiofur hydrochloride

Product identification

Medicine name:
CEFTIONIL

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. 8 day

- **Cattle**
  - 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

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Authorisation status:
Valid

Authorised in:
Italy

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:
26/11/2013

Manufacturing sites for batch release:
Laboratorios Maymo S.A.U.

Responsible authority:
Ministry Of Health

Authorisation number:
104629

Date of authorisation status change:
26/11/2013

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

Documents

Combined File of all Documents

English (PDF)

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Summary of Product Characteristics

Package Leaflet

Labelling

eu-PUAR-ceftionil-en.pdf