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Cefavex 50 mg/ml Suspension for Injection for Pigs and Cattle

Authorised

- Ceftiofur hydrochloride

Product identification

Medicine name:
Cefavex 50 mg/ml, suspension for injection for pigs and cattle
Cefavex 50 mg/ml Suspension for Injection for Pigs and Cattle

Active substance:
Ceftiofur hydrochloride

Target species:
Pig
Cattle

Route of administration:
Intramuscular use
Subcutaneous 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:

Intramuscular use:

- **Pig**
  - Meat and offal. 5 day

Subcutaneous use:

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

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:
Colourless glass type I vial of 100 ml, closed with grey coated bromobutyl rubber stopper and aluminium cap.Vial is individually packed in a carton box.One grouped as a clinical pack.
Colourless glass type I vial of 100 ml, closed with grey coated bromobutyl rubber stoppers and aluminium caps.Vials are individually packed in a carton box.Six vials are grouped as a clinical pack.
Colourless glass type I vial of 100 ml, closed with grey coated bromobutyl rubber stoppers and aluminium caps.Vials are individually packed in a carton box.Ten vials are grouped as a clinical pack.

Colourless glass type I vial of 100 ml, closed with grey coated bromobutyl rubber stoppers and aluminium caps.Vials are individually packed in a carton box.Twelve vials are grouped as a clinical pack.

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:
S P Veterinaria S.A.

Marketing authorisation date:
13/05/2013

Manufacturing sites for batch release:
S P Veterinaria S.A.

Responsible authority:
The Veterinary Medicines Directorate

Authorisation number:
Vm 36967/4002

Date of authorisation status change:
4/03/2024

Reference member state:
Ireland

Procedure number:
IE/V/0304/001

Concerned member states:
Bulgaria Germany Greece Hungary Italy Poland Portugal Romania 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

Combined File of all Documents