

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

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:

Italy

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:

12/06/2013

Manufacturing sites for batch release:

S P Veterinaria S.A.

Responsible authority:

Ministry Of Health

Authorisation number:

104519

Date of authorisation status change:

12/06/2013

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

English (PDF)

Published on: 22/12/2024

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Combined File of all Documents