Ceftiosan, 50 mg/ml, suspension for injection for pigs and cattle

Authorised

- Ceftiofur hydrochloride

Product identification

Medicine name:
Ceftiosan, 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. 8 day

Subcutaneous use:

- **Cattle**
  - Milk. no withdrawal period Should be 0 hours
  - 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:

Latvia

Package description:

Available only in Latvian

Carton box containing one glass vial, type II 100 ml, sealed with a bromobutyl rubber stopper and aluminium overseal.

Carton box containing one glass vial, type II 50 ml, sealed with a bromobutyl rubber stopper and aluminium overseal.

Polystyrene box containing 6 glass vials, type II 250 ml, sealed with bromobutyl rubber stopper and aluminium overseal.

Polystyrene box containing 15 glass vials, type II 50 ml, sealed with a bromobutyl rubber stopper and aluminium overseal.

Polystyrene box containing 12 glass vials, type II 100 ml, sealed with a bromobutyl rubber stopper and aluminium overseal.

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:
Alfasan Nederland B.V.

Marketing authorisation date:
19/01/2012

Manufacturing sites for batch release:
Alfasan Nederland B.V.

Responsible authority:
Food And Veterinary Service

Authorisation number:
V/DCP/12/0003

Date of authorisation status change:
19/01/2012

Reference member state:
Netherlands

Procedure number:
NL/V/0148/001

Concerned member states:
Austria Belgium Bulgaria Czechia Denmark Estonia France Germany Greece Hungary Ireland Italy Latvia Lithuania Poland Portugal Romania Slovenia 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: 16/06/2022
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Package Leaflet

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Labelling

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