Ceffect 25 mg/ml suspension for injection for cattle and pigs

Authorised

- Cefquinome sulfate

Product identification

Medicine name:
Ceffect 25 mg/ml suspension for injection for cattle and pigs
CEFFECT 25 mg/ml Injektionssuspension für Rinder und Schweine

Active substance:
Cefquinome sulfate

Target species:
Cattle
Pig

Route of administration:
Intramuscular use

Product details

Active substance and strength:
Cefquinome sulfate
29.64 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Suspension for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Cattle**
  - Meat and offal. 5 day
  - Milk. 24 hour
- **Pig**
  - Meat and offal. 3 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01DE90

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Austria

Available in:

Austria

Package description:

Carton with 1 colourless Type II glass vial containing 100 ml. Each vial is closed with a fluorinatedbromobutyl rubber stopper and sealed with an aluminium cap.
Carton with 6 colourless Type II glass vials containing 100 ml. Each vial is closed with a fluorinatedbromobutyl rubber stopper and sealed with an aluminium cap.
Carton with 12 colourless Type II glass vials containing 100 ml. Each vial is closed with a fluorinatedbromobutyl rubber stopper and sealed with an aluminium cap.
Carton with 1 colourless Type II glass vial containing 250 ml. Each vial is closed with a fluorinatedbromobutyl rubber stopper and sealed with an aluminium cap.

Carton with 6 colourless Type II glass vials containing 250 ml. Each vial is closed with a fluorinatedbromobutyl rubber stopper and sealed with an aluminium cap.

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:
Emdoka

Marketing authorisation date:
17/12/2012

Manufacturing sites for batch release:
Wirtschaftsgenossenschaft deutscher Tieraerzte eG

Responsible authority:
Austrian Agency For Health And Food Safety

Authorisation number:
8-01136

Date of authorisation status change:
17/12/2012

Reference member state:
Ireland

Procedure number:
IE/V/0470/001

Concerned member states:
Austria Belgium Czechia Germany Hungary Italy Luxembourg Netherlands Poland Portugal 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

English (PDF)
Published on: 26/10/2025
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Labelling

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Package Leaflet

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