

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

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:**

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Cattle

- Meat and offal. 5 day
- Milk. 24 hour

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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:

Germany

Available in:

Germany

Package description:

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.

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 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 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 1 colourless Type II glass vial containing 100 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:

27/11/2012

Manufacturing sites for batch release:

Wirtschaftsgenossenschaft deutscher Tieraerzte eG

Responsible authority:

Federal Office Of Consumer Protection And Food Safety

Authorisation number:

401649.00.00

Date of authorisation status change:

27/06/2018

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

Combined File of all Documents

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