

CEVAXEL 50 MG/ML POWDER AND SOLVENT FOR SOLUTION FOR INJECTION FOR CATTLE AND PIGS

Not
authorised

- Ceftiofur hydrochloride
- Water for injection

Product identification

Medicine name:

CEVAXEL 50 MG/ML POWDER AND SOLVENT FOR SOLUTION FOR INJECTION FOR CATTLE AND PIGS

Active substance:

Ceftiofur hydrochloride

Water for injection

Target species:

Cattle

Pig

Route of administration:

Intramuscular use

Product details

Active substance and strength:

Ceftiofur hydrochloride

53.50 milligram(s) / 1.00 millilitre(s)

Water for injection

1.00 millilitre(s) / 1.00 millilitre(s)

Pharmaceutical form:

Powder and solvent for solution for injection

Withdrawal period by route of administration:

Intramuscular use:

-

Cattle

- Meat and offal. 2 day

- Milk. 0 day

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Pig

- Meat and offal. 2 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01DD90

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Surrendered

Authorised in:

Estonia

Package description:

Box containing 1 vial of Cevaxel 50 mg/ml 1 g and box containing a vial of 20 ml solvent

Box containing 1 vial of Cevaxel 50 mg/ml, 4 g and box containing a vial of 80 ml solvent

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:

Ceva Sante Animale

Marketing authorisation date:

25/09/2008

Manufacturing sites for batch release:

Ceva Sante Animale

Responsible authority:

State Agency Of Medicines

Authorisation number:

1522

Date of authorisation status change:

27/06/2024

Reference member state:

France

Procedure number:

FR/V/0177/001

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Summary of Product Characteristics

This document does not exist in this language (English). You can find it in another language below.