

EFICUR 50 mg/ml suspension for injection for pigs and cattle

Authorised

- Ceftiofur hydrochloride

Product identification

Medicine name:

EFICUR 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 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01DD90

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Iceland

Package description:

Type II glass bottle of 50 ml. The bottles are closed with a Type I bromobutyl closure and aluminium cap. Pack size: Cardboard box with 1 glass bottle of 50 ml.

Type II glass bottle of 100 ml. The bottles are closed with a Type I bromobutyl closure and aluminium cap. Pack size: Cardboard box with 1 glass bottle of 100 ml.

Type II glass bottle of 250 ml. The bottles are closed with a Type I bromobutyl closure and aluminium cap. The 250 ml glass bottle has a colourless plastic package as a protective measure in order to avoid glass bottle breaking when it is being used. Pack size: Cardboard box with 1 glass bottle of 250 ml.

Type II glass bottles of 100 ml. The bottles are closed with a Type I bromobutyl closure and aluminium cap. Pack size: Cardboard box with 10 glass bottles of 100 ml.

Type II glass bottles of 100 ml. The bottles are closed with a Type I bromobutyl closure and aluminium capPack size: Cardboard box with 12 glass bottles of 100 ml.

Polyethylene terephthalate (PET) bottle of 50 ml. The bottles are closed with a Type I bromobutyl closure and aluminium capPack size: Cardboard box with 1 PET bottle of 50 ml.

Polyethylene terephthalate (PET) bottle of 100 ml. The bottles are closed with a Type I bromobutyl closure and aluminium capPack size: Cardboard box with 1 PET bottle of 100 ml.

Polyethylene terephthalate (PET) bottle of 250 ml. The bottles are closed with a Type I bromobutyl closure and aluminium capPack size: Cardboard box with 1 PET bottle of 250 ml

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:

Laboratorios Hipra S.A.

Marketing authorisation date:

29/01/2009

Manufacturing sites for batch release:

Laboratorios Hipra S.A.

Responsible authority:

Icelandic Medicines Agency

Authorisation number:

IS/2/08/013/01

Date of authorisation status change:

30/03/2012

Reference member state:

Ireland

Procedure number:

IE/V/0190/001

Concerned member states:

Austria Belgium Bulgaria Cyprus Czechia Denmark Estonia France
Germany Greece Hungary Iceland Italy Latvia Lithuania Luxembourg
Netherlands Poland Portugal Romania Slovakia 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: 28/09/2025

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Combined File of all Documents

Labelling

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Package Leaflet

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