

Fenflor 300 mg/ml solution for injection for cattle

Authorised

- Florfenicol

Product identification

Medicine name:

Fenflor 300 mg/ml solution for injection for cattle

Active substance:

Florfenicol

Target species:

Cattle

Route of administration:

Subcutaneous use
Intramuscular use

Product details

Active substance and strength:

Florfenicol
300.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:**Subcutaneous use:**

-

Cattle

- Meat and offal. 44 day by SC (at 40 mg/kg bodyweight, once)
- Milk. no withdrawal period

Not authorised for use in animals producing milk for human consumption.

Intramuscular use:

-

Cattle

- Meat and offal. 30 day by IM (at 20 mg/kg bodyweight, twice)
- Milk. no withdrawal period

Not authorised for use in animals producing milk for human consumption.

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01BA90

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Germany

Available in:

Germany

Package description:

(ID3) 250 millilitre(s): Box (board) with 1 Vial (brown glass) with 250 millilitre(s), closed with Stopfen (bromobutyl rubber) and Cap`` (aluminium)

(ID2) 100 millilitre(s): Box (board) with 1 Vial (brown glass) with 100 millilitre(s), closed with Stopfen (bromobutyl rubber) and Cap`` (aluminium)

(ID1) 50 millilitre(s): Box (board) with 1 Vial (brown glass) with 50 millilitre(s), closed with Stopfen (bromobutyl rubber) and Cap`` (aluminium)

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:

KRKA tovarna zdravil d.d. Novo mesto

Marketing authorisation date:

18/10/2010

Manufacturing sites for batch release:

KRKA tovarna zdravil d.d. Novo mesto

Responsible authority:

Federal Office Of Consumer Protection And Food Safety

Authorisation number:

401342.00.00

Date of authorisation status change:

21/10/2015

Reference member state:

Germany

Procedure number:

DE/V/0195/002

Concerned member states:

Austria Belgium Ireland Portugal Spain United Kingdom (Northern Ireland)

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

Documents

Combined File of all Documents

English (PDF)

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