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

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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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