

Fenflor 300 mg/ml solution for injection for pigs

Authorised

- Florfenicol

Product identification

Medicine name:

Fenflor 300 mg/ml solution for injection for pigs

Fenflor 300 mg/ml Oplossing voor injectie

Fenflor 300 mg/ml Solution injectable

Fenflor 300 mg/ml Injektionslösung

Active substance:

Florfenicol

Target species:

Pig

Route of administration:

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:**Intramuscular use:**

-

Pig

- Meat and offal. 18 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01BA90

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Belgium

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:

27/10/2008

Manufacturing sites for batch release:

KRKA tovarna zdravil d.d. Novo mesto

Responsible authority:

Federal Agency For Medicines And Health Products

Authorisation number:

BE-V327293

Date of authorisation status change:

27/10/2008

Reference member state:

Germany

Procedure number:

DE/V/0195/001

Concerned member states:

Austria Belgium Denmark Ireland Portugal 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

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

Package Leaflet

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

Combined File of all Documents

Labelling

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

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