

KEFLORIL 300 MG/ML SOLUTION FOR INJECTION FOR CATTLE AND PIGS

Authorised

- Florfenicol

Product identification

Medicine name:

KEFLORIL 300 MG/ML SOLUTION FOR INJECTION FOR CATTLE AND PIGS

Active substance:

Florfenicol

Target species:

Cattle

Pig

Route of administration:

Intramuscular use

Subcutaneous 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:**

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Cattle

- Milk. no withdrawal period

Milk: Not permitted for use in lactating animals producing milk for human consumption.

- Meat and offal. 30 day by IM (at 20 mg/kg bodyweight, twice): 30 days

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Pig

- Meat and offal. 18 day

Subcutaneous use:

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Cattle

- Meat and offal. 44 day by SC (at 40 mg/kg bodyweight, once): 44 days

- Milk. no withdrawal period

Milk: Not permitted for use in lactating 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:

France

Available in:

France

Package description:

Available only in [French](#)

Available only in [French](#)

Available only in [French](#)

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:

Vetoquinol S.A.

Marketing authorisation date:

20/07/2010

Manufacturing sites for batch release:

Vetoquinol S.A.

Responsible authority:

French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:

FR/V/2204461 7/2010

Date of authorisation status change:

16/06/2015

Reference member state:

France

Procedure number:

FR/V/0216/001

Concerned member states:

Austria Belgium Denmark Germany Ireland Italy Netherlands 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.

Combined File of all Documents

Package Leaflet and Labelling

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