

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

Kefloril 300 mg/ml, oplossing voor injectie voor runderen en varkens.

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:

Netherlands

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:

28/10/2020

Manufacturing sites for batch release:

Vetoquinol S.A.

Responsible authority:

Medicines Evaluation Board

Authorisation number:

REG NL 106149

Date of authorisation status change:

5/04/2022

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

Combined File of all Documents

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