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:

- **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

- **Pig**

  - Meat and offal. 18 day

Subcutaneous use:

- **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

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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

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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