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 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:

- **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:

Ireland

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 Ireland Limited

Marketing authorisation date:
24/09/2010

Manufacturing sites for batch release:
Vetoquinol

Responsible authority:
Health Products Regulatory Authority

Authorisation number:
VPA10983/053/001

Date of authorisation status change:
24/09/2010

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