MARBOCYL 100 MG/ML, SOLUTION FOR INJECTION FOR CATTLE AND PIGS

Authorised

- Marbofloxacin

Product identification

Medicine name:
MARBOCYL 100 MG/ML, SOLUTION FOR INJECTION FOR CATTLE AND PIGS

Active substance:
Marbofloxacin

Target species:
Cattle
Pig

Route of administration:
Intramuscular use
Subcutaneous use

Product details

Active substance and strength:
Marbofloxacin
100.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Cattle**

  - Meat and offal. 6 day

Indication : Respiratory Dosage : 2 m/kg for 3 to 5 days (IV/IM/SC)

  - Milk. 36 hour

Indication : Mastitis Dosage : 2mg/kg for 3 days (IV/IM/SC)

  - Meat and offal. 6 day

Indication : Mastitis Dosage : 2mg/kg for 3 days (IV/IM/SC)

  - Milk. 72 hour

Indication : Respiratory Dosage : 8 mg/kg on a single occasion (IM)

  - Meat and offal. 3 day

Indication : Respiratory Dosage : 8 mg/kg on a single occasion (IM)

  - Milk. 36 hour

Indication : Respiratory Dosage : 2 m/kg for 3 to 5 days (IV/IM/SC)

- **Pig**

  - Meat and offal. 4 day

Subcutaneous use:

- **Cattle**

  - Meat and offal. 6 day

Indication : Respiratory Dosage : 2 m/kg for 3 to 5 days (IV/IM/SC)

  - Milk. 36 hour

Indication : Mastitis Dosage : 2mg/kg for 3 days (IV/IM/SC)

- Meat and offal. 6 day

Indication : Mastitis Dosage : 2mg/kg for 3 days (IV/IM/SC)

- Milk. 72 hour

Indication : Respiratory Dosage : 8 mg/kg on a single occasion (IM)

- Meat and offal. 3 day

Indication : Respiratory Dosage : 8 mg/kg on a single occasion (IM)

- Milk. 36 hour

Indication : Respiratory Dosage : 2 m/kg for 3 to 5 days (IV/IM/SC)

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01MA93

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

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Authorisation status:
Valid

Authorised in:
Luxembourg

Available in:
Luxembourg

Package description:
Available only in French
Available only in French
Available only in French
Available only in French

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Vetoquinol S.A.

Marketing authorisation date:
3/11/1998

Manufacturing sites for batch release:
Vetoquinol Biowet Sp. z o.o.
Vetoquinol S.A.

Responsible authority:
Ministry Of Health And Social Security

Authorisation number:
V 808/98/08/0609

Date of authorisation status change:
3/11/1998

Reference member state:
France

Procedure number:
FR/V/0107/001

Concerned member states:
Austria Belgium Germany Greece Italy Luxembourg Portugal Spain

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Combined File of all Documents