MARBOCYL 20 MG/ML SOLUTION FOR INJECTION FOR CATTLE AND PIGS

Authorised

- Marbofloxacin

Product identification

Medicine name:
MARBOCYL 20 MG/ML SOLUTION FOR INJECTION FOR CATTLE AND PIGS

Active substance:
Marbofloxacin

Target species:
Cattle (calf)
Pig

Route of administration:
Intramuscular use
Subcutaneous use
Intravenous use

Product details

Active substance and strength:
Marbofloxacin

20.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:
Intramuscular use:

- **Cattle (calf)**
  - Meat and offal. 6 day
  - Milk. no withdrawal period

The veterinary medicinal product is not authorised for use in lactating animals producing milk for human consumption.

- **Pig**
  - Meat and offal. 4 day

Subcutaneous use:

- **Cattle (calf)**
  - Meat and offal. 6 day
  - Milk. no withdrawal period

The veterinary medicinal product is not authorised for use in lactating animals producing milk for human consumption.

Intravenous use:

- **Cattle (calf)**
  - Meat and offal. 6 day
  - Milk. no withdrawal period

The veterinary medicinal product is not authorised for use in lactating animals producing milk for human consumption.

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01MA93

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

Authorisation status:
Valid

Authorised in:
Portugal

Available in:
Portugal

Package description:
Box of one vial of 10 ml
Box of one vial of 250 ml
Box of one vial of 100 ml
Box of one vial of 50 ml
Box of one vial of 20 ml

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:
Vetoquinol S.A.

Marketing authorisation date:
29/03/1999

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

Responsible authority:

Directorate General For Food And Veterinary

Authorisation number:
51263

Date of authorisation status change:
4/04/2025

Reference member state:
France

Procedure number:
FR/V/0107/002

Concerned member states:
Austria Denmark 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

English (PDF)
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