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Meloxidyl 20 mg/ml - Solution for injection

Authorised

- Meloxicam

Product identification

Medicine name:
Meloxidyl 20 mg/ml - Solution for injection

Active substance:
Meloxicam

Target species:
Cattle
Horse
Pig

Route of administration:
Intramuscular use
Intravenous use
Subcutaneous use

Product details

Active substance and strength:
Meloxicam
20.00 milligram(s) / 1.00 Vial

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:
Intramuscular use:

- **Cattle**
  - Meat and offal. 15 day 15 days
  - Milk. 5 day 5 days
- **Horse**
  - Meat and offal. 5 day 5 days
- **Pig**
  - Meat and offal. 5 day 5 days

Intravenous use:

- **Cattle**
  - Meat and offal. 15 day 15 days
  - Milk. 5 day 5 days
- **Horse**
  - Meat and offal. 5 day 5 days
- **Pig**
  - Meat and offal. 5 day 5 days

Subcutaneous use:

- **Cattle**
  - Meat and offal. 15 day 15 days
  - Milk. 5 day 5 days
- **Horse**
  - Meat and offal. 5 day 5 days
- **Pig**
  - Meat and offal. 5 day 5 days

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QM01AC06

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Austria , Belgium , Bulgaria , Croatia , Cyprus , Czechia , Denmark , Estonia , Finland , France , Germany , Greece , Hungary , Iceland , Ireland , Italy , Latvia , Liechtenstein , Lithuania , Luxembourg , Malta , Netherlands , Norway , Poland , Portugal , Romania , Slovakia , Slovenia , Spain , Sweden , United Kingdom (Northern Ireland)

Available in:

Austria , Belgium , Bulgaria , Cyprus , Czechia , Denmark , Finland , France , Germany , Greece , Hungary , Ireland , Italy , Latvia , Luxembourg , Netherlands , Norway , Poland , Portugal , Slovakia , Slovenia , Spain , Sweden , United Kingdom (Northern Ireland)

Package description:

Packaging:Vial (glass), Package_size:1 vial, Content:50 ml

Packaging:Vial (glass), Package_size:1 vial, Content:100 ml

Packaging:Vial (glass), Package_size:1 vial, Content:250 ml

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:
Ceva Sante Animale

Marketing authorisation date:
15/01/2007

Manufacturing sites for batch release:
Ceva Sante Animale

Responsible authority:
European Commission

Authorisation number:
This information is not available for this product.

Date of authorisation status change:
26/03/2010

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

Documents

Combined File of all Documents

English (PDF)

Published on: 2/10/2026

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