Meloxidolor 5 mg/ml - Solution for injection

Authorised

- Meloxicam

Product identification

Medicine name:
Meloxidolor 5 mg/ml - Solution for injection

Active substance:
Meloxicam

Target species:
Cattle
Dog
Cat
Pig

Route of administration:
Intramuscular use
Intravenous use
Subcutaneous use

Product details

Active substance and strength:
Meloxicam

Presentation_strength:5 mg Reference:Ph.Eur Index:0

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Cattle**
  - Meat and offal. 15 day 15 days
- **Dog**
- **Cat**
- **Pig**
  - Meat and offal. 5 day 5 days

Intravenous use:

- **Cattle**
  - Meat and offal. 15 day 15 days
- **Dog**
- **Cat**
- **Pig**
  - Meat and offal. 5 day 5 days

Subcutaneous use:

- **Cattle**
  - Meat and offal. 15 day 15 days
- **Dog**
- **Cat**
- **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

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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)

Package description:
Packaging:multi-pack, Package_size:10 vials, Content:20 ml
Packaging:multi-pack, Package_size:5 vials, Content:20 ml
Packaging:Vial (glass), Package_size:1 vial, Content:100 ml
Packaging:Vial (glass), Package_size:1 vial, Content:20 ml
Packaging:Vial (glass), Package_size:1 vial, Content:10 ml

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Hybrid application (Article 13(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Le Vet Beheer B.V.

Marketing authorisation date:
22/04/2013

Manufacturing sites for batch release:
Produlab Pharma B.V.

Responsible authority:
European Commission

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

Date of authorisation status change:

22/04/2013

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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