Meloxidolor 40 mg/ml - Solution for injection

Not authorised

- Meloxicam

Product identification

Medicine name:
Meloxidolor 40 mg/ml - Solution for injection

Active substance:
Meloxicam

Target species:
Cattle
Horse

Route of administration:
Intravenous use

Product details

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

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:

Intravenous use:

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QM01AC06

Legal status of supply:

Medicinal product subject to medical prescription

Authorisation status:

Surrendered

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)

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:
3/07/2015

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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