File downloaded on 2026-11-07
Source URL: https://medicines.health.europa.eu/veterinary/en/600000985709

Melovem 15 mg/ml - Oral suspension (Horse)

Authorised

- Meloxicam

Product identification

Medicine name:
Melovem 15 mg/ml - Oral suspension (Horse)

Active substance:
Meloxicam

Target species:
Horse

Route of administration:
Oral use

Product details

Active substance and strength:
Meloxicam
15.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Oral suspension

Withdrawal period by route of administration:

Oral use:

- **Horse**
  - Meat and offal. 3 day 3 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 , Czechia , Denmark , Estonia , Finland , France , Germany , Hungary , Italy , Latvia , Lithuania , Luxembourg , Netherlands , Norway , Poland , Portugal , Romania , Slovakia , Spain , Sweden

Package description:

Packaging:Bottle (HDPE), Package_size:1 bottle, Content:250 ml

Packaging:Bottle (HDPE), Package_size:1 bottle, Content:500 ml

Packaging:Bottle (HDPE), Package_size:1 Bottle, Content:100 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:
Dopharma Research B.V.

Marketing authorisation date:
7/07/2009

Manufacturing sites for batch release:
Dopharma B.V.

Responsible authority:
European Commission

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

Date of authorisation status change:
26/06/2023

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: 16/01/2024
Updated on: 14/03/2026
Download

ema-puar-v0152-melovem-vra-0015-en.pdf