File downloaded on 2026-11-08
Source URL: https://medicines.health.europa.eu/veterinary/en/600000107125

Meloxoral 1 mg - Chewable tablet

Authorised

- Meloxicam

Product identification

Medicine name:
Meloxoral 1 mg - Chewable tablet

Active substance:
Meloxicam

Target species:
Dog

Route of administration:
Oral use

Product details

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

Pharmaceutical form:
Chewable tablet

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:
Germany

Package description:
Packaging:Blister (OPA/Alu/PVC), Package_size:30 tablets
Packaging:Blister (OPA/Alu/PVC), Package_size:50 tablets
Packaging:Blister (OPA/Alu/PVC), Package_size:100 tablets

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:
Dechra Regulatory B.V.

Marketing authorisation date:
19/11/2010

Manufacturing sites for batch release:
Produlab Pharma B.V.
Lelypharma B.V.
Genera Analitika d.o.o.

Responsible authority:

European Commission

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

Date of authorisation status change:
30/11/2022

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: 14/03/2026
Download

ema-puar-meloxoral-v-000151-var-x-0015-en.pdf

ema-puar-meloxoral-v-151-par-en.pdf