Loxitab 2.5 mg - Tablet

Authorised

- Meloxicam

Product identification

Medicine name:
Loxitab 2.5 mg - Tablet

Active substance:
Meloxicam

Target species:
Dog

Route of administration:
Oral use

Product details

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

Pharmaceutical form:
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:
Belgium , Estonia , Germany , Greece , Ireland , Italy , Latvia , Lithuania , Netherlands , Portugal , Spain

Package description:
Packaging:Blister (Alu/PVC/PE/PVDC), Package_size:5 blisters x 10 tablets
Packaging:Blister (Alu/PVC/PE/PVDC), Package_size:10 blisters x 10 tablets
Packaging:Blister (Alu/PVC/PE/PVDC), Package_size:3 blisters x 10 tablets
Packaging:Blister (Alu/PVC/PE/PVDC), Package_size:1 blister x 10 tablets

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Hybrid application – change in strength (Article 19(1)(a) of Regulation (EU) 2019/6)

Marketing authorisation holder:
CP-Pharma Handelsgesellschaft mbH

Marketing authorisation date:
19/10/2023

Manufacturing sites for batch release:
CP-Pharma Handelsgesellschaft mbH

Responsible authority:

European Commission

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

Date of authorisation status change:
19/10/2023

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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