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Coxatab 25 mg - Chewable tablet

Authorised

- Firocoxib
- Firocoxib

Product identification

Medicine name:
Coxatab 25 mg - Chewable tablet

Active substance:
Firocoxib
Firocoxib

Target species:
Dog
Dog

Route of administration:
Oral use
Oral use

Product details

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

25.00 milligram(s) / 1.00 Tablet

Pharmaceutical form:

Chewable tablet

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QM01AH90

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:

Denmark , Finland , Germany , Greece , Norway , Spain , Sweden

Package description:

Packaging:Blister (Alu/PVC/PE/PVDC), Package_size:10 tablets
Packaging:Blister (Alu/PVC/PE/PVDC), Package_size:200 tablets
Packaging:Blister (Alu/PVC/PE/PVDC), Package_size:30 tablets
Packaging:Blister (Alu/PVC/PE/PVDC), Package_size:100 tablets
Packaging:Blister (Alu/PVC/PE/PVDC), Package_size:20 tablets
Packaging:Blister (Alu/PVC/PE/PVDC), Package_size:50 tablets
Packaging:Blister (Alu/PVC/PE/PVDC), Package_size:10 tablets
Packaging:Blister (Alu/PVC/PE/PVDC), Package_size:200 tablets
Packaging:Blister (Alu/PVC/PE/PVDC), Package_size:30 tablets
Packaging:Blister (Alu/PVC/PE/PVDC), Package_size:100 tablets
Packaging:Blister (Alu/PVC/PE/PVDC), Package_size:20 tablets
Packaging:Blister (Alu/PVC/PE/PVDC), Package_size:50 tablets

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:
CP-Pharma Handelsgesellschaft mbH

Marketing authorisation date:
12/08/2022

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:
12/08/2022

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: 13/03/2026

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