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Previcox 227 mg - Chewable tablet

Authorised

- Firocoxib

Product identification

Medicine name:
Previcox 227 mg - Chewable tablet

Active substance:
Firocoxib

Target species:
Dog

Route of administration:
Oral use

Product details

Active substance and strength:
Firocoxib
227.28 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:
Austria , Belgium , Czechia , Estonia , France , Latvia , Lithuania , Luxembourg , Poland , Slovakia , Spain

Package description:
Packaging:Blister (Alu/PVC), Package_size:10 tablets
Packaging:Blister (Alu/PVC), Package_size:30 tablets
Packaging:Blister (Alu/PVC), Package_size:180 tablets
Packaging:Bottle (PET), Package_size:60 tablets

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Complete application (stand-alone) - Council Directive 81/851/EEC

Marketing authorisation holder:
Boehringer Ingelheim Vetmedica GmbH

Marketing authorisation date:
13/09/2004

Manufacturing sites for batch release:

Boehringer Ingelheim Animal Health France

Responsible authority:
European Commission

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

Date of authorisation status change:
13/09/2004

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
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