

# Daxocox 15 mg - Tablet

Authorised

- Enflicoxib

## Product identification

**Medicine name:**

Daxocox 15 mg - Tablet

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**Active substance:**

Enflicoxib

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**Target species:**

Dog

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**Route of administration:**

Oral use

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## Product details

**Active substance and strength:**

Enflicoxib

15.00 milligram(s) / 1.00 Tablet

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**Pharmaceutical form:**

Tablet

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QM01AH95

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**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)

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**Available in:**

Austria , Belgium , Bulgaria , Croatia , Cyprus , Czechia , Denmark , Germany , Greece , Hungary , Ireland , Italy , Malta , Netherlands , Norway , Poland , Portugal , Romania , Slovakia , Spain

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**Package description:**

Packaging:Blister (Alu), Package\_size:12 (3 x 4) tablets  
Packaging:Blister (Alu), Package\_size:20 (4 x 5) tablets  
Packaging:Blister (Alu), Package\_size:24 (6 x 4) tablets  
Packaging:Blister (Alu), Package\_size:100 (10 x 10) tablets  
Packaging:Blister (Alu), Package\_size:10 (1 x 10) tablets  
Packaging:Blister (Alu), Package\_size:50 (5 x 10) tablets  
Packaging:Blister (Alu), Package\_size:4 (1 x 4) tablets  
Packaging:Blister (Alu), Package\_size:5 (1 x 5) tablets

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Full application - New active substance (Article 12(3) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Ecuphar NV

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**Marketing authorisation date:**

20/04/2021

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**Manufacturing sites for batch release:**

Lelypharma B.V.

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**Responsible authority:**

European Commission

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**Authorisation number:**

This information is not available for this product.

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**Date of authorisation status change:**

19/12/2024

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To consult adverse reactions on veterinary medicinal products please go to [www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Combined File of all Documents

English (PDF)

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