

# Onsior 10 mg - Tablet (dogs)

Authorised

- Robenacoxib

## Product identification

**Medicine name:**

Onsior 10 mg - Tablet (dogs)

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**Active substance:**

Robenacoxib

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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:**

Robenacoxib

Presentation\_strength:10 mg Reference:Monograph Index:0

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**Pharmaceutical form:**

Tablet

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

QM01AH91

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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 , Cyprus , Czechia , Denmark , Estonia , Finland , France , Germany , Greece , Hungary , Ireland , Italy , Latvia , Lithuania , Luxembourg , Netherlands , Norway , Poland , Portugal , Romania , Slovakia , Slovenia , Spain , Sweden , United Kingdom (Northern Ireland)

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**Package description:**

Packaging:Blister (Alu/Alu), Package\_size:28 tablets

Packaging:Blister (Alu/Alu), Package\_size:70 tablets

Packaging:Blister (Alu/Alu), Package\_size:7 tablets

Packaging:Blister (Alu/Alu), Package\_size:14 tablets

Packaging:Blister (Alu/Alu), Package\_size:30 x 1 tablets

Packaging:Blister (Alu/Alu), Package\_size:60 x 1 tablets

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Full application (Article 12(3) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Elanco GmbH

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**Marketing authorisation date:**

16/12/2008

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**Manufacturing sites for batch release:**

Elanco France S.A.S.

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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:**

11/12/2020

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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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