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# Onsior 10 mg - Tablet (dogs)

Authorised

- Robenacoxib

## Product identification

**Medicine name:**
Onsior 10 mg - Tablet (dogs)

**Active substance:**
Robenacoxib

**Target species:**
Dog

**Route of administration:**
Oral use

## Product details

**Active substance and strength:**
Robenacoxib

Presentation_strength:10 mg Reference:Monograph Index:0

**Pharmaceutical form:**
Tablet

**Anatomical therapeutic chemical veterinary (ATCvet) codes:**
QM01AH91

**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 , 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)

**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

## Additional information

**Entitlement type:**
Marketing Authorisation

**Legal basis of product authorisation:**
Full application (Article 12(3) of Directive No 2001/82/EC)

**Marketing authorisation holder:**
Elanco GmbH

**Marketing authorisation date:**
16/12/2008

**Manufacturing sites for batch release:**
Elanco France S.A.S.

**Responsible authority:**
European Commission

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

**Date of authorisation status change:**
16/12/2008

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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ema-puar-onsior-v-127-var-ii-24-g-en.pdf

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