

Onsior 20 mg - Tablet (dogs)

Authorised

- Robenacoxib

Product identification

Medicine name:

Onsior 20 mg - Tablet (dogs)

Active substance:

Robenacoxib

Target species:

Dog

Route of administration:

Oral use

Product details

Active substance and strength:

Robenacoxib

Presentation_strength:20 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:60 x 1 tablets

Packaging:Blister (Alu/Alu), Package_size:30 x 1 tablets

Packaging:Blister (Alu/Alu), Package_size:70 tablets

Packaging:Blister (Alu/Alu), Package_size:28 tablets

Packaging:Blister (Alu/Alu), Package_size:14 tablets

Packaging:Blister (Alu/Alu), Package_size:7 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:

11/12/2020

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: 2/05/2024

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