

Loxicom 0.5 mg/ml - Oral suspension (cats)

Authorised

- Meloxicam

Product identification

Medicine name:

Loxicom 0.5 mg/ml - Oral suspension (cats)

Active substance:

Meloxicam

Target species:

Cat

Route of administration:

Oral use

Product details

Active substance and strength:

Meloxicam

0.50 milligram(s) / 1.00 Bottle

Pharmaceutical form:

Oral suspension

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QM01AC06

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 , Greece , Hungary , Ireland , Latvia , Lithuania , Luxembourg , Malta , Netherlands , Poland , Portugal , Romania , Slovenia , Spain , Sweden , United Kingdom (Northern Ireland)

Package description:

Packaging:Bottle (PET), measuring syringe (PP/PE), Package_size:1 bottle with dosing syringe, Content:30 ml

Packaging:Bottle (PET), measuring syringe (PP/PE), Package_size:1 bottle with dosing syringe, Content:15 ml

Packaging:Bottle (PET), measuring syringe (PP/PE), Package_size:1 bottle with dosing syringe, Content:5 ml

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:

Norbrook Laboratories (Ireland) Limited

Marketing authorisation date:

10/02/2009

Manufacturing sites for batch release:

Norbrook Laboratories Limited
Norbrook Manufacturing Limited

Responsible authority:

European Commission

Authorisation number:

This information is not available for this product.

Date of authorisation status change:

23/03/2010

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: 6/11/2025

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