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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 , Italy , 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:5 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:30 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: 5/11/2026
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