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Meloxidyl 0.5 mg/ml - Oral suspension

Authorised

- Meloxicam
- Meloxicam

Product identification

Medicine name:
Meloxidyl 0.5 mg/ml - Oral suspension

Active substance:
Meloxicam
Meloxicam

Target species:
Cat
Cat

Route of administration:
Oral use
Oral use

Product details

Active substance and strength:
Meloxicam
0.50 milligram(s) / 1.00 Bottle

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

Package description:
Packaging:Bottle (HDPE) with measuring syringe (LDPE), Package_size:1 bottle + 1 syringe, Content:15 ml
Packaging:Bottle (glass) with measuring syringe (LDPE), Package_size:1 bottle + 1 syringe, Content:5 ml
Packaging:Bottle (HDPE) with measuring syringe (LDPE), Package_size:1 bottle + 1 syringe, Content:15 ml
Packaging:Bottle (HDPE) with measuring syringe (LDPE), Package_size:1 bottle + 1 syringe, Content:4 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:
Ceva Sante Animale

Marketing authorisation date:
15/01/2007

Manufacturing sites for batch release:
Ceva Sante Animale

Responsible authority:
European Commission

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

Date of authorisation status change:
26/08/2010

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: 2/10/2026
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