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Rheumocam 1.5 mg/ml - Oral suspension

Authorised

- Meloxicam

Product identification

Medicine name:
Rheumocam 1.5 mg/ml - Oral suspension

Active substance:
Meloxicam

Target species:
Dog

Route of administration:
Oral use

Product details

Active substance and strength:
Meloxicam
1.50 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)

Package description:
Packaging:Bottle (PET), syringe (PP), Package_size:1 bottle + 1 syringe, Content:42 ml
Packaging:Bottle (PET), syringe (PP), Package_size:1 bottle + 1 syringe, Content:100 ml
Packaging:Bottle (PET), syringe (PP), Package_size:1 bottle + 1 syringe, Content:200 ml
Packaging:Bottle (HDPE), syringe (PP), Package_size:1 bottle + 1 syringe, Content:15 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:
Chanelle Pharmaceuticals Manufacturing Limited

Marketing authorisation date:
10/01/2008

Manufacturing sites for batch release:
Chanelle Pharmaceuticals Manufacturing Limited

Responsible authority:
European Commission

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

Date of authorisation status change:
10/01/2008

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: 13/03/2026
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