

Loxicom 5 mg/ml - Solution for injection (cats, dogs)

Authorised

- Meloxicam

Product identification

Medicine name:

Loxicom 5 mg/ml - Solution for injection (cats, dogs)

Active substance:

Meloxicam

Target species:

Dog

Cat

Route of administration:

Intravenous use

Subcutaneous use

Product details

Active substance and strength:

Meloxicam

Presentation_strength:5 mg Reference:B P Index:0

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:**Intravenous use:**

- Dog
- Cat

Subcutaneous use:

- Dog
- Cat

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:Vial (glass), Package_size:1 vial, Content:100 ml

Packaging:Vial (glass), Package_size:1 vial, Content:20 ml

Packaging:Vial (glass), Package_size:1 vial, Content:10 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:

10/02/2009

To consult adverse reactions on veterinary medicinal products please go to
www.adrreports.eu/vet

Documents

Combined File of all Documents

English (PDF)

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