Metaxx 1.5 mg/ml oral suspension for dogs

Authorised

- Meloxicam

Product identification

Medicine name:
Metaxx 1.5 mg/ml oral suspension for dogs
Metaxx 1,5 mg/ml perorálna suspenzia pre psy

Active substance:
Meloxicam

Target species:
Dog

Route of administration:
Oral use

Product details

Active substance and strength:
Meloxicam
1.50 milligram(s) / 1.00 millilitre(s)

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:
Slovakia

Package description:
5 ml (in a 10 ml sized bottle). Cardboard box with one HDPE bottle with an LDPE syringe adapter and closed with a polypropylene screw cap. Polypropylene measuring syringe of 3 mL.
10 ml. Cardboard box with one HDPE bottle with an LDPE syringe adapter and closed with a polypropylene screw cap. Polypropylene measuring syringe of 3 mL.
25 ml. Cardboard box with one HDPE bottle with an LDPE syringe adapter and closed with a polypropylene screw cap. Polypropylene measuring syringe of 3 mL.
50 ml. Cardboard box with one HDPE bottle with an LDPE syringe adapter and closed with a polypropylene screw cap. Polypropylene measuring syringe of 3 mL.
125 ml. Cardboard box with one HDPE bottle with an LDPE syringe adapter and closed with a polypropylene screw cap. Polypropylene measuring syringe of 3 mL.

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Generic application (Article 18 of Regulation (EU) 2019/6)

Marketing authorisation holder:
Alfasan Nederland B.V.

Marketing authorisation date:
20/01/2025

Manufacturing sites for batch release:
Alfasan Nederland B.V.
Produlab Pharma B.V.

Responsible authority:
Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:
96/054/DC/24-S

Date of authorisation status change:
20/01/2025

Reference member state:
Netherlands

Procedure number:
NL/V/0399/001

Concerned member states:
Austria Belgium Bulgaria Croatia Cyprus Czechia Denmark Estonia Finland France Germany Greece Hungary Iceland Ireland Italy Latvia Lithuania Luxembourg Norway Poland Portugal Romania Slovakia Slovenia Spain Sweden United Kingdom (Northern Ireland)

Generic of:
600000001530

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

Documents

Combined File of all Documents

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