

Metaxx 1.5 mg/ml oral suspension for dogs

Authorised

- Meloxicam

Product identification

Medicine name:

Metaxx 1.5 mg/ml oral suspension for dogs

Mexxam Vet 1,5 mg/ml Suspension zum Eingeben für Hunde

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:

Austria

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:

1/07/2024

Manufacturing sites for batch release:

Alfasan Nederland B.V.
Produlab Pharma B.V.

Responsible authority:

Austrian Agency For Health And Food Safety

Authorisation number:

842208

Date of authorisation status change:

1/07/2024

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

Package Leaflet

This document does not exist in this language (English). You can find it in another language below.

Summary of Product Characteristics

This document does not exist in this language (English). You can find it in another language below.

Labelling

This document does not exist in this language (English). You can find it in another language below.