

Animeloxan 1.5 mg/ml oral suspension for dogs

Authorised

- Meloxicam

Product identification

Medicine name:

Animeloxan 1.5 mg/ml oral suspension for dogs

Animeloxan 1,5 mg/ml sospensione orale per cani

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:

Italy

Package description:

(ID3) 100 millilitre(s): Box (Cardboard) with 1 Bottle (Polyethylen hoher Dichte) with 100 millilitre(s), closed with Lid and Verschluss mit Tropfer (Polyethylen, Polyethylen)
(ID1) 10 millilitre(s): Box (Cardboard) with 1 Bottle (Polyethylen hoher Dichte) with 10 millilitre(s), closed with Lid and Verschluss mit Tropfer (Polyethylen, Polyethylen)
(ID6) 125 millilitre(s): Box (Cardboard) with 1 Bottle (Polyethylen hoher Dichte) with 125 millilitre(s), closed with Verschluss mit Tropfer and Lid (Polyethylen, Polyethylen)
(ID4) 25 millilitre(s): Box (Cardboard) with 1 Bottle (Polyethylen hoher Dichte) with 25 millilitre(s), closed with Verschluss mit Tropfer and Lid (Polyethylen, Polyethylen)
(ID5) 50 millilitre(s): Box (Cardboard) with 1 Bottle (Polyethylen hoher Dichte) with 50 millilitre(s), closed with Lid and Verschluss mit Tropfer (Polyethylen, Polyethylen)

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:

aniMedica GmbH

Marketing authorisation date:

1/08/2024

Manufacturing sites for batch release:

aniMedica GmbH

Responsible authority:

Ministry Of Health

Authorisation number:

105827

Date of authorisation status change:

1/08/2024

Reference member state:

Germany

Procedure number:

DE/V/0310/001

Concerned member states:

Austria Denmark Hungary Italy Netherlands Poland Spain

United Kingdom (Northern Ireland)

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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