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 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

Withdrawal period by route of administration:

Oral use:

- **Dog**

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:
(ID3) 100 millilitre(s): Box (Cardboard) with 1 Bottle (High Density PolyEthylene) with 100 millilitre(s), closed with Lid and Dropper applicator (PolyEthylene, PolyEthylene)
(ID1) 10 millilitre(s): Box (Cardboard) with 1 Bottle (High Density PolyEthylene) with 10 millilitre(s), closed with Lid and Dropper applicator (PolyEthylene, PolyEthylene)
(ID6) 125 millilitre(s): Box (Cardboard) with 1 Bottle (High Density PolyEthylene) with 125 millilitre(s), closed with Dropper applicator and Lid (PolyEthylene, PolyEthylene)
(ID4) 25 millilitre(s): Box (Cardboard) with 1 Bottle (High Density PolyEthylene) with 25 millilitre(s), closed with Dropper applicator and Lid (PolyEthylene, PolyEthylene)
(ID5) 50 millilitre(s): Box (Cardboard) with 1 Bottle (High Density PolyEthylene) with 50 millilitre(s), closed with Lid and Dropper applicator (PolyEthylene, PolyEthylene)

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:
2/07/2008

Manufacturing sites for batch release:
aniMedica GmbH

Responsible authority:
Austrian Agency For Health And Food Safety

Authorisation number:
8-00750

Date of authorisation status change:
2/07/2008

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

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.

Combined File of all Documents
