Acticam 1 mg chewable tablets for Dogs

Not authorised

- Meloxicam

Product identification

Medicine name:
Acticam 1 mg chewable tablets for Dogs
Acticam, 1.0mg, Žvýkací tableta

Active substance:
Meloxicam

Target species:
Dog

Route of administration:
Oral use

Product details

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

Pharmaceutical form:
Chewable tablet

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:
Surrendered

Authorised in:
Czechia

Package description:
10 tablets (10x1) in a carton box. PVC/PVDC-Alu blister pack made of clear PVC/PVDC and Plain Alu foil. Each blister has 10 tablets.
100 tablets (10x10) in a carton box. PVC/PVDC-Alu blister pack made of clear PVC/PVDC and Plain Alu foil. Each blister has 10 tablets.
20 tablets (10x2) in a carton box. PVC/PVDC-Alu blister pack made of clear PVC/PVDC and Plain Alu foil. Each blister has 10 tablets.
500 tablets (10x50) in a carton box. PVC/PVDC-Alu blister pack made of clear PVC/PVDC and Plain Alu foil. Each blister has 10 tablets.

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:
Ecuphar

Marketing authorisation date:

19/06/2012

Manufacturing sites for batch release:
Accord Healthcare Limited
Ecuphar

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

Authorisation number:
96/065/12-C

Date of authorisation status change:
21/12/2021

Reference member state:
Netherlands

Procedure number:
NL/V/0134/001

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

---