Acticam 2.5 mg chewable tablets for Dogs

Authorised

Meloxicam

Product identification

Medicine name:

Acticam 2.5 mg chewable tablets for Dogs Acticam, 2.5 mg, Žvýkací tableta

Active substance:

Meloxicam

Target species:

Dog

Route of administration:

Oral use

Product details

Active substance and strength:

Meloxicam

2.50 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:

Valid

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.

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.

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.

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

Responsible authority:

Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:

96/066/12-C

Date of authorisation status change:

19/06/2012

Reference member state:

Netherlands

Procedure number:

NL/V/0134/002

Concerned member states:

Czechia Germany Hungary Italy Luxembourg Portugal Slovakia Spain

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

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