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Acticam (WD) Expired - 1.5 mg/ml - Oral suspension

Not authorised

- Meloxicam

Product identification

Medicine name:
Acticam (WD) Expired - 1.5 mg/ml - Oral suspension

Active substance:
Meloxicam

Target species:
Dog

Route of administration:
Oral use

Product details

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

Pharmaceutical form:
Oral suspension

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QM01AC06

Legal status of supply:
Medicinal product subject to medical prescription

Authorisation status:
Surrendered

Authorised in:
Austria , Belgium , Bulgaria , Croatia , Cyprus , Czechia , Denmark , Estonia , Finland , France , Germany , Greece , Hungary , Iceland , Ireland , Italy , Latvia , Liechtenstein , Lithuania , Luxembourg , Malta , Netherlands , Norway , Poland , Portugal , Romania , Slovakia , Slovenia , Spain , Sweden , United Kingdom (Northern Ireland)

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:
9/12/2008

Manufacturing sites for batch release:
Ecuphar

Responsible authority:
European Commission

Authorisation number:
This information is not available for this product.

Date of authorisation status change:

9/12/2008

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