Intercam 0.5 mg/ml oral suspension for cats

Authorised

- Meloxicam

Product identification

Medicine name:
Intercam 0.5 mg/ml oral suspension for cats

Active substance:
Meloxicam

Target species:
Cat

Route of administration:
Oral use

Product details

Active substance and strength:
Meloxicam
0.50 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:
Ireland

Package description:
Polypropylene bottle containing 5 ml with a tamper proof child resistant closures.Each bottle is packed in a cardboard box with a 1 ml measuring syringe (barrel in polypropylene and plunger/piston in high density polyethylene).
Polypropylene bottle containing 3 ml with a tamper proof child resistant closures.Each bottle is packed in a cardboard box with a 1 ml measuring syringe (barrel in polypropylene and plunger/piston in high density polyethylene).
White high density polyethylene bottle containing 10 ml with a tamper proof child resistant closure.Each bottle is packed in a cardboard box with a 1 ml measuring syringe (barrel in polypropylene and plunger/piston in high density polyethylene).
White high density polyethylene bottle containing 15 ml with a tamper proof child resistant closure.Each bottle is packed in a cardboard box with a 1 ml measuring syringe (barrel in polypropylene and plunger/piston in high density 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:
C&H Generics Limited

Marketing authorisation date:

15/07/2022

Manufacturing sites for batch release:
Chanelle Pharmaceuticals Manufacturing Limited

Responsible authority:
Health Products Regulatory Authority

Authorisation number:
VPA23184/001/001

Date of authorisation status change:
15/07/2022

Reference member state:
Ireland

Procedure number:
IE/V/0882/001

Concerned member states:
France Portugal Spain

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

Documents

Summary of Product Characteristics