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Inflacam 15 mg/ml - Oral suspension

Authorised

- Meloxicam

Product identification

Medicine name:
Inflacam 15 mg/ml - Oral suspension

Active substance:
Meloxicam

Target species:
Horse

Route of administration:
Oral use

Product details

Active substance and strength:
Meloxicam
1.50 percent / 1.00 Bottle

Pharmaceutical form:
Oral suspension

Withdrawal period by route of administration:

Oral use:

- **Horse**
  - Meat and offal. 3 day 3 days

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QM01AC06

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

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)

Package description:

Packaging:Bottle (HDPE), syringe (PP), Package_size:1 bottle + 1 syringe, Content:250 ml

Packaging:Bottle (HDPE), syringe (PP), Package_size:1 bottle + 1 syringe, Content:100 ml

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:

Chanelle Pharmaceuticals Manufacturing Limited

Marketing authorisation date:
9/12/2011

Manufacturing sites for batch release:
Chanelle Pharmaceuticals Manufacturing Limited

Responsible authority:
European Commission

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

Date of authorisation status change:
9/12/2011

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

Documents

Combined File of all Documents

English (PDF)
Published on: 22/09/2026
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