

Rheumocam 15 mg/ml - Oral suspension (horses)

Authorised

- Meloxicam

Product identification

Medicine name:

Rheumocam 15 mg/ml - Oral suspension (horses)

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: 100 ml

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

Content: 250 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 Ltd

Marketing authorisation date:

10/01/2008

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:

24/01/2011

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

Documents

Combined File of all Documents

English (PDF)

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