

# Emdocam 5 mg/ml - Solution for injection (cattle, pigs)

Authorised

- Meloxicam

## Product identification

**Medicine name:**

Emdocam 5 mg/ml - Solution for injection (cattle, pigs)

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**Active substance:**

Meloxicam

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**Target species:**

Cattle

Pig

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**Route of administration:**

Intramuscular use

Intravenous use

Subcutaneous use

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## Product details

**Active substance and strength:**

Meloxicam

5.00 milligram(s) / 1.00 Vial

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**Pharmaceutical form:**

Solution for injection

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**Withdrawal period by route of administration:****Intramuscular use:**

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**Cattle**

- Meat and offal. 15 day 15 days

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**Pig**

- Meat and offal. 5 day 5 days

**Intravenous use:**

- 

**Cattle**

- Meat and offal. 15 day 15 days

- 

**Pig**

- Meat and offal. 5 day 5 days

**Subcutaneous use:**

- 

**Cattle**

- Meat and offal. 15 day 15 days

- 

**Pig**

- Meat and offal. 5 day 5 days

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QM01AC06

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**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)

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**Package description:**

Packaging:Vial (glass), Package\_size:1 vial, Content:250 ml

Packaging:Vial (glass), Package\_size:1 vial, Content:100 ml

Packaging:Vial (glass), Package\_size:1 vial, Content:50 ml

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Generic application (Article 13(1) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Emdoka

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**Marketing authorisation date:**

18/08/2011

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**Manufacturing sites for batch release:**

Produlab Pharma B.V.

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**Responsible authority:**

European Commission

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**Authorisation number:**

This information is not available for this product.

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**Date of authorisation status change:**

26/04/2021

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To consult adverse reactions on veterinary medicinal products please go to [www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Combined File of all Documents

English (PDF)

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