

# EMDOFLUXIN 50 MG/ML SOLUTION FOR INJECTION FOR CATTLE, HORSES AND PIGS

Authorised

- Flunixin meglumine

## Product identification

**Medicine name:**

EMDOFLUXIN 50 MG/ML SOLUTION FOR INJECTION FOR CATTLE, HORSES AND PIGS

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**Active substance:**

Flunixin meglumine

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**Target species:**

Cattle

Pig

Equid

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**Route of administration:**

Intramuscular use

Intravenous use

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## Product details

**Active substance and strength:**

Flunixin meglumine

83.00 milligram(s) / 1.00 millilitre(s)

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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. 31 day
- Milk. 36 hour

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**Pig**

- Meat and offal. 24 day

**Intravenous use:**

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**Cattle**

- Meat and offal. 4 day
- Milk. 24 hour

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**Equid**

- Meat and offal. 5 day
- Milk. no withdrawal period

Milk: Not authorised for use in mares producing milk for human consumption.

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

QM01AG90

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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:**

Spain

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**Package description:**

Available only in [French](#)

Available only in [French](#)

Available only in [French](#)

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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:**

16/09/2020

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**Manufacturing sites for batch release:**

Produlab Pharma B.V.

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**Responsible authority:**

Spanish Agency Of Medicines And Medical Devices

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**Authorisation number:**

3938 ESP

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**Date of authorisation status change:**

1/01/2023

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**Reference member state:**

France

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**Procedure number:**

FR/V/0417/001

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**Concerned member states:**

Austria Belgium Bulgaria Croatia Denmark Estonia Germany Hungary  
Ireland Italy Latvia Lithuania Luxembourg Netherlands Slovenia Spain  
United Kingdom (Northern Ireland)

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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

### Package Leaflet

This document does not exist in this language (English). You can find it in another language below.

### Labelling

This document does not exist in this language (English). You can find it in another language below.

### Summary of Product Characteristics

This document does not exist in this language (English). You can find it in another language below.

Combined File of all Documents

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