

# MEGANYL 50 mg/ml Solution for Injection for cattle, pigs and horses.

Authorised

- Flunixin meglumine

## Product identification

**Medicine name:**

MEGANYL 50 mg/ml Solution for Injection for cattle, pigs and horses.

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**Active substance:**

Flunixin meglumine

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**Target species:**

Cattle

Horse

Pig

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**Route of administration:**

Intravenous use

Intramuscular 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:**

**Intravenous use:**

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**Cattle**

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

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**Horse**

- Meat and offal. 4 day
- Milk. no withdrawal period

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

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**Cattle**

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

•

**Horse**

- Meat and offal. 4 day
- Milk. no withdrawal period

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

**Intramuscular use:**

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**Pig**

- Meat and offal. 24 day

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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:**

Bulgaria

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**Available in:**

Bulgaria

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**Package description:**

box containing 1 vial of 250 ml

box containing 1 vial of 100 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:**

Laboratorios Syva S.A.

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**Marketing authorisation date:**

7/08/2019

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**Manufacturing sites for batch release:**

Laboratorios Syva S.A.

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**Responsible authority:**

Bulgarian Food Safety Authority

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**Authorisation number:**

0022-2907

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**Date of authorisation status change:**

7/08/2019

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**Reference member state:**

Spain

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**Procedure number:**

ES/V/0249/001

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**Concerned member states:**

Belgium Bulgaria Cyprus France Greece Hungary Italy Poland Portugal  
Romania

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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

### Summary of Product Characteristics

English (PDF)

Published on: 5/04/2023

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### Package Leaflet

English (PDF)

Published on: 30/10/2025

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Labelling

English (PDF)

Published on: 23/03/2023

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