

# Vetaflumex 50 mg/ml, solução injectável para bovinos, equinos e suínos

Authorised

- Flunixin meglumine
- Flunixin meglumine

## Product identification

### Medicine name:

Vetaflumex 50 mg/ml, solução injectável para bovinos, equinos e suínos

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### Active substance:

Flunixin meglumine

Flunixin meglumine

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### Target species:

Pig

Cattle

Horse

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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)

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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#### **Pig**

- Meat and offal. 20 day

#### **Intravenous use:**

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#### **Cattle**

- Meat and offal. 4 day

- Milk. 1 day

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#### **Horse**

- Meat and offal. 1 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:**

Lithuania

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**Available in:**

Lithuania

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**Package description:**

Polypropylene bottles closed with a rubber stopper oversealed with aluminium caps with a capacity of 50 ml.

Polypropylene bottles closed with a rubber stopper oversealed with aluminium caps with a capacity of 100 ml.

Polypropylene bottles closed with a rubber stopper oversealed with aluminium caps with a capacity of 250 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:**

Przedsiębiorstwo Wielobranzowe Vet-Agro Sp. z o.o.

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**Marketing authorisation date:**

26/11/2014

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**Manufacturing sites for batch release:**

Vet-Agro Trading Sp. z o.o.

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**Responsible authority:**

State Food And Veterinary Service

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**Authorisation number:**

LT/2/14/2263/001-003

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**Date of authorisation status change:**

26/01/2020

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**Reference member state:**

Portugal

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**Procedure number:**

PT/V/0116/001

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**Concerned member states:**

Bulgaria Greece Hungary Lithuania Romania Slovakia

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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

Combined File of all Documents

English (PDF)

Published on: 1/08/2025

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