

# Finadyne, 50 mg/ml, solution for injection

Authorised

- Flunixin meglumine

## Product identification

**Medicine name:**

Finadyne, 50 mg/ml, solution for injection

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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

Intramuscular 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. 5 day
- Milk. no withdrawal period

Leche: Su uso no está autorizado en animales cuya leche se utiliza para el consumo humano.

**Intramuscular use:**

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**Pig**

- Meat and offal. 24 day

**Intramuscular use:**

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**Cattle**

- Meat and offal. 31 day

**Intramuscular use:**

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**Cattle**

- Milk. 36 hour

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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:**

Poland

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**Package description:**

1 Cardboard box with 10 x 50 ml vial  
1 Cardboard box with 1 x 250 ml vial  
1 Cardboard box with 1 x 100 ml vial  
1 Cardboard box with 1 x 50 ml vial  
1 Cardboard box with 6 x 50 ml vial  
1 Cardboard box with 6 x 250 ml vial  
1 Cardboard box with 10 x 100 ml vial

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Intervet International B.V.

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**Marketing authorisation date:**

17/05/2000

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**Manufacturing sites for batch release:**

Trirx Segre

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**Responsible authority:**

Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

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**Authorisation number:**

**Date of authorisation status change:**

17/05/2000

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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.

### Summary of Product Characteristics

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.

### Combined File of all Documents