

# 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 est? 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:**

France

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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

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**Marketing authorisation date:**

4/08/1981

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**Manufacturing sites for batch release:**

Trirx Segre

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**Responsible authority:**

French Agency For Food, Environmental And Occupational Health & Safety

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**Authorisation number:**

FR/V/0046543 8/1981

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**Date of authorisation status change:**

4/08/2011

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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

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

Package Leaflet and Labelling

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

Combined File of all Documents

Package Leaflet