

# LIVERFINE 100 mg/ml SOLUCION INYECTABLE

Authorised

- Fibric acid

## Product identification

**Medicine name:**

LIVERFINE 100 mg/ml SOLUCION INYECTABLE

**Active substance:**

Fibric acid

**Target species:**

Cattle

Goat

Horse

Pig

Dog

**Route of administration:**

Intramuscular use

Intraperitoneal use

Intravenous use

## Product details

**Active substance and strength:**

Fibric acid

100.00 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. 0 day
- Milk. 0 hour

- 

**Goat**

- Meat and offal. 0 day
- Milk. 0 hour

- 

**Horse**

- Meat and offal. 0 day

- 

**Pig**

- Meat and offal. 0 day

**Intraperitoneal use:**

- 

**Cattle**

- Meat and offal. 0 day
- Milk. 0 hour

- 

**Goat**

- Meat and offal. 0 day
- Milk. 0 hour

- 

**Horse**

- Meat and offal. 0 day

- 

**Pig**

- Meat and offal. 0 day

**Intravenous use:**

- 

**Cattle**

- Meat and offal. 0 day

- Milk. 0 hour

- 

**Goat**

- Meat and offal. 0 day

- Milk. 0 hour

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**Horse**

- Meat and offal. 0 day

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**Pig**

- Meat and offal. 0 day

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

QA05BA

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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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**Available in:**

Spain

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**Package description:**

Available only in [Spanish](#)

Available only in [Spanish](#)

Available only in [Spanish](#)

Available only in [Spanish](#)

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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:**

Fatro S.p.A.

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**Marketing authorisation date:**

10/09/2008

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**Manufacturing sites for batch release:**

Fatro S.p.A.

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**Responsible authority:**

Spanish Agency For Medicines And Medical Devices

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**Authorisation number:**

1918 ESP

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**Date of authorisation status change:**

10/09/2008

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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

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.