

# Tolfine

Authorised

- Tolfenamic acid

## Product identification

**Medicine name:**

Tolfine

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**Active substance:**

Tolfenamic acid

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**Target species:**

Cattle

Pig

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**Route of administration:**

Intramuscular use

Intravenous use

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## Product details

**Active substance and strength:**

Tolfenamic acid

40.00 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. 12 day

- Milk. 0 hour

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**Pig**

- Meat and offal. 16 day

**Intravenous use:**

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**Cattle**

- Meat and offal. 4 day

- Milk. 24 hour

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

QM01AG02

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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:**

Ireland

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**Available in:**

Ireland

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**Package description:**

The product Tolfine is packaged in an amber type I glass vial of 50 ml. The vial is closed with a chlorobutyl rubber stopper with aluminium flip cap. Each vial is packaged in a cardboard box.

The product Tolfine is packaged in an amber type I glass vial of 100 ml. The vial is closed with a chlorobutyl rubber stopper with aluminium flip cap. Each vial is packaged in a cardboard box.

The product Tolfine is packaged in an amber type I glass vial of 250 ml. The vial is closed with a chlorobutyl rubber stopper with aluminium flip cap. Each vial is packaged in a cardboard box.

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## Additional information

### **Entitlement type:**

Marketing Authorisation

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### **Legal basis of product authorisation:**

Full application (Article 12(3) of Directive No 2001/82/EC)

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### **Marketing authorisation holder:**

Vetoquinol Ireland Limited

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### **Marketing authorisation date:**

4/02/2000

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### **Manufacturing sites for batch release:**

Vetoquinol S.A.

Vetoquinol Biowet Sp. z o.o.

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### **Responsible authority:**

Health Products Regulatory Authority

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### **Authorisation number:**

VPA10983/031/001

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### **Date of authorisation status change:**

4/02/2000

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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

Summary of Product Characteristics