

# QUINOBAL 100 mg/ml SOLUCION INYECTABLE PARA BOVINO PORCINO OVINO Y CAPRINO

Authorised

- Enrofloxacin

## Product identification

### Medicine name:

QUINOBAL 100 mg/ml SOLUCION INYECTABLE PARA BOVINO PORCINO OVINO Y CAPRINO

### Active substance:

Enrofloxacin

### Target species:

Cattle

Sheep

Goat

Pig

### Route of administration:

Subcutaneous use

Intravenous use

Intramuscular use

## Product details

### Active substance and strength:

Enrofloxacin

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:

#### Subcutaneous use:

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##### Cattle

- Meat and offal. no withdrawal period Carne: 5 días (IV); 12 días (SC)

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##### Sheep

- Meat and offal. 4 day

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##### Goat

- Meat and offal. 6 day

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##### Cattle

- Milk. no withdrawal period Leche: 3 días (IV); 4 días (SC)

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##### Sheep

- Milk. 3 day

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##### Goat

- Milk. 4 day

#### Intravenous use:

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#### **Cattle**

- Meat and offal. no withdrawal period Carne: 5 días (IV); 12 días (SC)

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#### **Cattle**

- Milk. no withdrawal period Leche: 3 días (IV); 4 días (SC)

#### **Intramuscular use:**

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#### **Pig**

- Meat and offal. 13 day

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

QJ01MA90

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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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#### **Package description:**

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:**

Global Vet Health S.L.

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**Marketing authorisation date:**

16/04/2019

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**Manufacturing sites for batch release:**

S P Veterinaria S.A.

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**Responsible authority:**

Spanish Agency Of Medicines And Medical Devices

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**Authorisation number:**

3776 ESP

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**Date of authorisation status change:**

16/04/2019

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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.

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