

# DORAFLOX 100 mg/ml solution for injection for cattle and pigs

Authorised

- Enrofloxacin

## Product identification

**Medicine name:**

DORAFLOX 100 mg/ml solution for injection for cattle and pigs

---

**Active substance:**

Enrofloxacin

---

**Target species:**

Pig

Cattle

---

**Route of administration:**

Intramuscular use

Intravenous use

Subcutaneous use

---

## Product details

**Active substance and strength:**

Enrofloxacin

100.00 milligram(s) / 1.00 millilitre(s)

---

**Pharmaceutical form:**

Solution for injection

---

**Withdrawal period by route of administration:****Intramuscular use:**

- 

**Pig**

- Meat and offal. 13 day

**Intravenous use:**

- 

**Cattle**

- Meat and offal. 5 day

- Milk. 3 day

**Subcutaneous use:**

- 

**Cattle**

- Meat and offal. 12 day

- Milk. 4 day

---

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

QJ01MA90

---

**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

---

**Authorisation status:**

Valid

---

**Authorised in:**

Germany

---

**Available in:**

Germany

---

**Package description:**

Type II Amber glass vial of 250 ml closed with bromobutyl rubber stopper and aluminium seal. One vial of 250 ml is available in a cardboard box.

Type II Amber glass vial of 100 ml closed with bromobutyl rubber stopper and aluminium seal. One vial of 100 ml is available in a cardboard box.

---

## Additional information

**Entitlement type:**

Marketing Authorisation

---

**Legal basis of product authorisation:**

Generic application (Article 13(1) of Directive No 2001/82/EC)

---

**Marketing authorisation holder:**

Dopharma Research B.V.

---

**Marketing authorisation date:**

31/03/2011

---

**Manufacturing sites for batch release:**

Dopharma B.V.

---

**Responsible authority:**

Federal Office Of Consumer Protection And Food Safety

---

**Authorisation number:**

401391.02.01

---

**Date of authorisation status change:**

17/06/2016

---

**Reference member state:**

Ireland

---

**Procedure number:**

IE/V/0261/001

---

**Concerned member states:**

Belgium Germany Greece Hungary Netherlands Romania

United Kingdom (Northern Ireland)

---

To consult adverse reactions on veterinary medicinal products please go to [www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Combined File of all Documents

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