

# Enrox Max 100 mg/ml Injektionslösung für Rinder und Schweine

Authorised

- Enrofloxacin

## Product identification

**Medicine name:**

Enrox Max 100 mg/ml Injektionslösung für Rinder und Schweine

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**Active substance:**

Enrofloxacin

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**Target species:**

Pig

Cattle

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**Route of administration:**

Intramuscular use

Intravenous use

Subcutaneous use

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## 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:**

**Intramuscular use:**

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**Pig**

- Meat and offal. 12 day

7,5 mg/kg KGW/Tag, ev. 2. Injektion in Abstand von 48 Std.

**Intravenous use:**

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**Cattle**

- Meat and offal. 7 day 5 mg/kg KGW/Tag

- Milk. 72 hour 5 mg/kg KGW/Tag

**Subcutaneous use:**

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**Cattle**

- Meat and offal. 14 day

7,5 mg/kg KGW/Tag, ev. 2. Injektion in Abstand von 48 Std.

- Milk. 120 hour 7,5 mg/kg KGW/Tag, ev. 2. Injektion in Abstand von 48 Std.

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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:**

Bulgaria

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**Available in:**

Bulgaria

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**Package description:**

Cardboard box with one amber glass multi-dose vial (Type II) containing 100 ml with bromobutyl rubber stopper and aluminium seal.

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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:**

KRKA tovarna zdravil d.d. Novo mesto

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**Marketing authorisation date:**

10/09/2013

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**Manufacturing sites for batch release:**

KRKA tovarna zdravil d.d. Novo mesto

TAD Pharma GmbH

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**Responsible authority:**

Bulgarian Food Safety Authority

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**Authorisation number:**

0022-2102

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**Date of authorisation status change:**

10/09/2013

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**Reference member state:**

Austria

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**Procedure number:**

AT/V/0010/001

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**Concerned member states:**

Belgium Bulgaria Czechia France Germany Hungary Italy Latvia Lithuania  
Portugal Romania Slovakia Slovenia Spain United Kingdom (Northern Ireland)

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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

English (PDF)

Published on: 29/12/2023

Updated on: 13/03/2026

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### Package Leaflet and Labelling

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