

# Enroxal 100 mg/ml oral solution for chickens and turkeys

Authorised

- Enrofloxacin

## Product identification

**Medicine name:**

Enroxal 100 mg/ml oral solution for chickens and turkeys

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**Active substance:**

Enrofloxacin

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**Target species:**

Turkey

Chicken

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**Route of administration:**

In drinking water 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:**

Oral solution

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**Withdrawal period by route of administration:**

**In drinking water use:**

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**Turkey**

- Meat and offal. 13 day
- Egg. no withdrawal period

Not authorised for use in birds producing eggs for human consumption. Do not administer to layer replacement birds within 14 days of coming into lay.

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**Chicken**

- Egg. no withdrawal period

Not authorised for use in birds producing eggs for human consumption. Do not administer to layer replacement birds within 14 days of coming into lay.

- Meat and offal. 7 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:**

Germany

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**Available in:**

Germany

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**Package description:**

(ID3) 5 litre(s): unspecified outer container with 1 Bottle (high-density polyethylene) with 5 litre(s)

(ID2) 1 litre(s): unspecified outer container with 1 Bottle (high-density polyethylene) with 1 litre(s)

(ID1) 100 millilitre(s): unspecified outer container with 1 Bottle (brown glass) with 100 millilitre(s)

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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:**

TAD Pharma GmbH

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**Marketing authorisation date:**

25/03/2013

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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:**

Federal Office Of Consumer Protection And Food Safety

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**Authorisation number:**

401913.00.00

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**Date of authorisation status change:**

17/02/2018

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**Reference member state:**

Germany

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**Procedure number:**

DE/V/0336/001

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**Concerned member states:**

Belgium Bulgaria Cyprus Italy Netherlands

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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

Combined File of all Documents

English (PDF)

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