

# Lanflox 100 mg/ml Solution for Use in Drinking Water for Chickens and Turkeys

Authorised

- Enrofloxacin

## Product identification

**Medicine name:**

LANFLOX 100 mg/ml Solution for use in drinking water for Chickens and Turkeys  
Lanflox 100 mg/ml Solution for Use in Drinking Water 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:**

Solution for use in drinking water

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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

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**Chicken**

- Egg. no withdrawal period
- 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:**

United Kingdom (Northern Ireland)

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**Available in:**

United Kingdom (Northern Ireland)

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**Package description:**

(ID3) 5 litre(s): unspecified outer container with 1 Barrel (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) 250 millilitre(s): unspecified outer container with 1 Bottle (High Density PolyEthylene) with 250 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:**

Vetpharma Animal Health S.L.

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**Marketing authorisation date:**

12/03/2010

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**Manufacturing sites for batch release:**

Laboratorios Karizoo S.A.

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**Responsible authority:**

The Veterinary Medicines Directorate

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**Authorisation number:**

Vm 32509/4002

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**Date of authorisation status change:**

2/09/2024

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**Reference member state:**

Germany

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**Procedure number:**

DE/V/0346/001

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**Concerned member states:**

Belgium France Hungary Ireland Netherlands Romania

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

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**Source URL:** <https://medicines.health.europa.eu/veterinary/600000047513>