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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

Active substance:
Enrofloxacin

Target species:
Turkey
Chicken

Route of administration:
In drinking water use

Product details

Active substance and strength:
Enrofloxacin
100.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for use in drinking water

Withdrawal period by route of administration:
In drinking water use:

- **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.

- **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

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01MA90

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
United Kingdom (Northern Ireland)

Available in:
United Kingdom (Northern Ireland)

Package description:
(ID1) 250 millilitre(s): unspecified outer container with 1 Bottle (high-density polyethylene) with 250 millilitre(s)

(ID2) 1 litre(s): unspecified outer container with 1 Bottle (high-density polyethylene) with 1 litre(s)
(ID3) 5 litre(s): unspecified outer container with 1 Barrel (high-density polyethylene) with 5 litre(s)

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:
Vetpharma Animal Health S.L.

Marketing authorisation date:
12/03/2010

Manufacturing sites for batch release:
Laboratorios Karizoo S.A.

Responsible authority:
The Veterinary Medicines Directorate

Authorisation number:
Vm 32509/4002

Date of authorisation status change:
2/09/2024

Reference member state:
Germany

Procedure number:
DE/V/0346/001

Concerned member states:
Belgium France Hungary Ireland Netherlands Romania

United Kingdom (Northern Ireland)

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Summary of Product Characteristics

Package Leaflet