Enrotron 100 mg/ml Solution for Use in Drinking Water for Chicken, Turkeys and Rabbits

Authorised

- Enrofloxacin

Product identification

Medicine name:
Enrotron 100 mg/ml Solution for Use in Drinking Water for Chicken, Turkeys and Rabbits

Active substance:
Enrofloxacin

Target species:
Turkey
Rabbit
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
- **Rabbit**
  - Meat and offal. 15 day
- **Chicken**
  - 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:
Cardboard box containing 6 x 1000 ml opaque polyethylene bottle with opaque polyethylene tamper evident screw cap and polypropylene measuring pitcher.
Cardboard box containing 12 x 100 ml opaque polyethylene bottle with opaque polyethylene tamper evident screw cap and polypropylene measuring pitcher.

1000 ml opaque polyethylene bottle with opaque polyethylene tamper evident screw cap and polypropylene measuring pitcher
Cardboard box containing 100 ml opaque polyethylene bottle with opaque polyethylene tamper evident screw cap and polypropylene measuring pitcher.

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:
aniMedica GmbH

Marketing authorisation date:
20/06/2013

Manufacturing sites for batch release:
aniMedica GmbH
Industrial Veterinaria S.A.

Responsible authority:
The Veterinary Medicines Directorate

Authorisation number:
Vm 24745/4022

Date of authorisation status change:
18/06/2024

Reference member state:
Netherlands

Procedure number:
NL/V/0165/003

Concerned member states:

Cyprus France Germany Ireland Poland United Kingdom (Northern Ireland)

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