

Baytril 100 mg/ml solution for use in drinking water for chickens, turkeys and rabbits

Authorised

- Enrofloxacin

Product identification

Medicine name:

Baytril 100 mg/ml solution for use in drinking water for chickens, turkeys and rabbits

Active substance:

Enrofloxacin

Target species:

Turkey
Chicken
Rabbit

Route of administration:

Oral use

Product details

Active substance and strength:

Enrofloxacin
100.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Oral solution

Withdrawal period by route of administration:**Oral use:**

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Turkey

- Meat and offal. 13 day

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Chicken

- Meat and offal. 7 day

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Rabbit

- Meat and offal. 3 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01MA90

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Ireland

Available in:

Ireland

Package description:

5,000 ml HDPE canister with an aluminium / HDPE screw closure. The containers are provided with a graduated polypropylene measuring cup.

1,000 ml high density polyethylene (HDPE) bottles with an HDPE insert and a polypropylene screw closure. The containers are provided with a graduated polypropylene measuring cup.

500 ml high density polyethylene (HDPE) bottles with an HDPE insert and a polypropylene screw closure. The containers are provided with a graduated polypropylene measuring cup.

100 ml high density polyethylene (HDPE) bottles with an HDPE insert and a polypropylene screw closure. The containers are provided with a graduated polypropylene measuring cup.

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:

Elanco GmbH

Marketing authorisation date:

1/10/1988

Manufacturing sites for batch release:

KVP Pharma+Veterinaer Produkte GmbH

Responsible authority:

Health Products Regulatory Authority

Authorisation number:

VPA22020/063/001

Date of authorisation status change:

1/10/1988

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

Documents

Summary of Product Characteristics

Combined File of all Documents