

Enrotron 100 mg/ml oral solution for chicken and turkeys

Authorised

- Enrofloxacin

Product identification

Medicine name:

Enrotron 100 mg/ml oral solution for chicken and turkeys

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:

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Turkey

- Meat and offal. 13 day

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Rabbit

- Meat and offal. 15 day

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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:

Poland

Available in:

Poland

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:

13/12/2013

Manufacturing sites for batch release:

aniMedica GmbH

Industrial Veterinaria S.A.

Responsible authority:

Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

Authorisation number:

2312

Date of authorisation status change:

13/12/2013

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

Documents

Labelling

This document does not exist in this language (English). You can find it in another language below.

Summary of Product Characteristics

This document does not exist in this language (English). You can find it in another language below.

Package Leaflet

This document does not exist in this language (English). You can find it in another language below.