Enroxal 100 mg/ml oral solution for chickens and turkeys

Authorised

- Enrofloxacin

Product identification

Medicine name:
Enroxal 100 mg/ml oral solution 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:
Oral solution

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:
Bulgaria

Available in:
Bulgaria

Package description:
(ID3) 5 litre(s): unspecified outer container with 1 Bottle (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) 100 millilitre(s): unspecified outer container with 1 Bottle (brown glass) with 100 millilitre(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:
KRKA tovarna zdravil d.d. Novo mesto

Marketing authorisation date:
12/03/2013

Manufacturing sites for batch release:
KRKA tovarna zdravil d.d. Novo mesto
TAD Pharma GmbH

Responsible authority:
Bulgarian Food Safety Authority

Authorisation number:
0022-1972

Date of authorisation status change:
12/03/2013

Reference member state:
Germany

Procedure number:
DE/V/0336/001

Concerned member states:
Belgium Bulgaria Cyprus Italy Netherlands

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

Documents

Summary of Product Characteristics

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

Combined File of all Documents

Package Leaflet

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

Labelling

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