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

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Authorisation status:
Valid

Authorised in:
Italy

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:
4/02/2014

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

Responsible authority:
Ministry Of Health

Authorisation number:
104590

Date of authorisation status change:
4/02/2014

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

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Combined File of all Documents

English (PDF)
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