ENROFLOXAN, 100mg/ml, Perorální roztok

Authorised

- Enrofloxacin

Product identification

Medicine name:
ENROFLOXAN, 100mg/ml, Perorální roztok

Active substance:
Enrofloxacin

Target species:
Chicken (broiler)
Cattle (pre-ruminant)
Pig

Route of administration:
In drinking water/milk 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/milk use:

- **Chicken (broiler)**
  - Meat and offal. 7 day
  - Egg. no withdrawal period

Nepoužívat u kuřic během 14 dní před počátkem snášky.,

- **Cattle (pre-ruminant)**
  - Meat and offal. 12 day

- **Pig**
  - Meat and offal. 12 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01MA90

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Czechia

Package description:
Available only in Czech
Available only in Czech

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:
Ing. Vojtech Lorenc Csc.

Marketing authorisation date:
6/08/2012

Manufacturing sites for batch release:
Biofaktor Sp. z o.o.

Responsible authority:
Institute For State Control Of Veterinary Biologicals And Medicines

Authorisation number:
96/088/12-C

Date of authorisation status change:
6/08/2012

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

Documents

Summary of Product Characteristics

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Package Leaflet

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

Labelling

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