Enroxil 50 mg/ml šķīdums injekcijām teļiem, cūkām, aitām, kazām un suņiem

Authorised

- Enrofloxacin

Product identification

Medicine name:
Enroxil 50 mg/ml šķīdums injekcijām teļiem, cūkām, aitām, kazām un suņiem

Active substance:
Enrofloxacin

Target species:
Pig
Cattle (calf)
Goat
Sheep
Dog

Route of administration:
Intramuscular use
Intravenous use
Subcutaneous use

Product details

Active substance and strength:
Enrofloxacin
50.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Pig**
  - Meat and offal. 13 day

Intravenous use:

- **Cattle (calf)**
  - Meat and offal. 5 day

Subcutaneous use:

- **Goat**
  - Meat and offal. 6 day
  - Milk. 4 day
- **Sheep**
  - Milk. 3 day
  - Meat and offal. 4 day
- **Cattle (calf)**
  - 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:
Latvia

Available in:
Latvia

Package description:
Available only in Latvian
Available only in Latvian

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:
30/11/1995

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

Responsible authority:
Food And Veterinary Service

Authorisation number:

V/NRP/95/0195

Date of authorisation status change:
30/11/1995

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

Documents

Combined File of all Documents

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