Enrox Max 100 mg/ml Injektionslösung für Rinder und Schweine

Authorised

- Enrofloxacin

Product identification

Medicine name:
Enrox Max 100 mg/ml Injektionslösung für Rinder und Schweine

Active substance:
Enrofloxacin

Target species:
Pig
Cattle

Route of administration:
Intramuscular use
Intravenous use
Subcutaneous use

Product details

Active substance and strength:
Enrofloxacin

100.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:
Intramuscular use:

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

7,5 mg/kg KGW/Tag, ev. 2. Injektion in Abstand von 48 Std.

Intravenous use:

- **Cattle**
  - Meat and offal. 7 day 5 mg/kg KGW/Tag
  - Milk. 72 hour 5 mg/kg KGW/Tag

Subcutaneous use:

- **Cattle**
  - Meat and offal. 14 day

7,5 mg/kg KGW/Tag, ev. 2. Injektion in Abstand von 48 Std.

  - Milk. 120 hour 7,5 mg/kg KGW/Tag, ev. 2. Injektion in Abstand von 48 Std.

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01MA90

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

Authorisation status:
Valid

Authorised in:

Hungary

Available in:
Hungary

Package description:
Cardboard box with one amber glass multi-dose vial (Type II) containing 100 ml with bromobutyl rubber stopper and aluminium seal.

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:
25/09/2013

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

Responsible authority:
Directorate Of Veterinary Medicinal Products

Authorisation number:
This information is not available for this product.

Date of authorisation status change:
25/09/2013

Reference member state:
Austria

Procedure number:

AT/V/0010/001

Concerned member states:
Belgium Bulgaria Czechia France Germany Hungary Italy Latvia Lithuania Portugal Romania Slovakia Slovenia Spain United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics
at-puar-atv0010001-mr-enroex-en.pdf