

Enroxil, 100mg/ml, Injekční roztok

Authorised

- Enrofloxacin

Product identification

Medicine name:

Enroxil, 100mg/ml, Injekční roztok

Active substance:

Enrofloxacin

Target species:

Cattle

Pig

Route of administration:

Intravenous use

Subcutaneous use

Intramuscular 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:

Intravenous use:

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Cattle

- Meat and offal. 5 day
- Milk. 3 day

Subcutaneous use:

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Cattle

- Meat and offal. 12 day
- Milk. 4 day

Intramuscular use:

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Pig

- Meat and offal. 13 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

Available in:

Czechia

Package description:

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:

KRKA tovarna zdravil d.d. Novo mesto

Marketing authorisation date:

27/07/1995

Manufacturing sites for batch release:

KRKA tovarna zdravil d.d. Novo mesto

Responsible authority:

Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:

96/677/95-C

Date of authorisation status change:

27/07/1995

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.

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.