

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:

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Pig

- Meat and offal. 12 day

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

Intravenous use:

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Cattle

- Meat and offal. 7 day 5 mg/kg KGW/Tag

- Milk. 72 hour 5 mg/kg KGW/Tag

Subcutaneous use:

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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:

Austria

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:

24/09/2013

Manufacturing sites for batch release:

KRKA tovarna zdravil d.d. Novo mesto

TAD Pharma GmbH

Responsible authority:

Austrian Agency For Health And Food Safety

Authorisation number:

835160

Date of authorisation status change:

24/09/2013

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

English (PDF)

Published on: 29/12/2023

Updated on: 8/04/2026

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Package Leaflet

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Labelling

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