

Baytril 100 mg/ml raztopina za injiciranje

Authorised

- Enrofloxacin

Product identification

Medicine name:

Baytril 100 mg/ml raztopina za injiciranje

Active substance:

Enrofloxacin

Target species:

Cattle

Sheep

Goat

Pig

Route of administration:

Subcutaneous use

Intravenous 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:

Subcutaneous use:

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Cattle

- Meat and offal. 5 day

Govedo: Po intravenskem injiciranju: Meso in organi: 5 dni. Mleko: 3 dni.

- Milk. 3 day

Govedo: Po intravenskem injiciranju: Meso in organi: 5 dni. Mleko: 3 dni.

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Sheep

- Meat and offal. 4 day Ovce: Meso in organi: 4 dni. Mleko: 3 dni.

- Milk. 3 day Ovce: Meso in organi: 4 dni. Mleko: 3 dni.

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Goat

- Meat and offal. 6 day Koze: Meso in organi: 6 dni. Mleko: 4 dni.

- Milk. 4 day Koze: Meso in organi: 6 dni. Mleko: 4 dni.

Intravenous use:

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Cattle

- Meat and offal. 5 day

Po intravenskem injiciranju: Meso in organi: 5 dni. Mleko: 3 dni.

- Milk. 3 day Po intravenskem injiciranju: Meso in organi: 5 dni. Mleko: 3 dni.

Intramuscular use:

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Pig

- Meat and offal. 13 day Prašiči: Meso in organi: 13 dni.

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01MA90

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Slovenia

Available in:

Slovenia

Package description:

Available only in Slovenian

Available only in Slovenian

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:

Elanco Animal Health GmbH

Marketing authorisation date:

4/01/2005

Manufacturing sites for batch release:

KVP Pharma+Veterinaer Produkte GmbH

Responsible authority:

Agency For Medicinal Products And Medical Devices Of The Republic Of Slovenia

Authorisation number:

NP/V/0030/002

Date of authorisation status change:

4/01/2005

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.