**Source URL:** https://medicines.health.europa.eu/veterinary/en/600000095726

# BAYTRIL 50mg/ml ενέσιμο διάλυμα

Authorised

Enrofloxacin

## **Product identification**

#### **Medicine name:**

BAYTRIL 50mg/ml ενέσιμο διάλυμα

#### **Active substance:**

Enrofloxacin

## **Target species:**

Cattle (calf)

Dog

Goat

Sheep

Cat

Pig

### **Route of administration:**

Intravenous use

Intramuscular 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: Intravenous use:

•

#### Cattle (calf)

- Meat and offal. 5 day
- Meat and offal. 12 day

•

#### Goat

- Meat and offal. 6 day
- Milk. 4 day

•

## Sheep

- Meat and offal. 4 day
- Milk. 3 day

•

## Pig

- Meat and offal. 13 day

#### Intramuscular use:

•

## Cattle (calf)

- Meat and offal. 5 day
- Meat and offal. 12 day

## Goat

- Meat and offal. 6 day
- Milk. 4 day

#### •

## Sheep

- Meat and offal. 4 day
- Milk. 3 day

#### •

## Pig

- Meat and offal. 13 day

#### **Subcutaneous use:**

#### •

## Cattle (calf)

- Meat and offal. 5 day
- Meat and offal. 12 day

#### •

### Goat

- Meat and offal. 6 day
- Milk. 4 day

#### •

## Sheep

- Meat and offal. 4 day
- Milk. 3 day

## Pig

- Meat and offal. 13 day

## Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01MA90

Authorisation status:	
Valid	
<b>Authorised in:</b> Cyprus	
<b>Available in:</b> Cyprus	
Package description:	
Available only in <u>Greek</u>	
Available only in <u>Greek</u>	
Additional information	
Entitlement type: Marketing Authorisation	
Legal basis of product autho	risation:
Full application - Known active s	ubstance (Article 12(3) of Directive No 2001/82/EC)
Marketing authorisation hold	ler:
Elanco Animal Health GmbH	

## Manufacturing sites for batch release:

KVP Pharma+Veterinaer Produkte GmbH

## **Responsible authority:**

Veterinary Services, Ministry Of Agriculture, Natural Resources And Environment

#### **Authorisation number:**

12631

2/02/1990

## **Date of authorisation status change:**

27/11/2012

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

## **Documents**

Combined File of all Documents

This document does not exist in this language (English). You can find it in another language below.