BELAMOX, milteliai ir tirpiklis injekciniam tirpalui

Authorised

- Amoxicillin sodium

Product identification

Medicine name:
BELAMOX, milteliai ir tirpiklis injekciniam tirpalui

Active substance:
Amoxicillin sodium

Target species:
Cattle (calf)
Horse
Pig
Cattle

Route of administration:
Intramuscular use
Intravenous use
Subcutaneous use

Product details

Active substance and strength:
Amoxicillin sodium

5.30 gram(s) / 1.00 Vial

Pharmaceutical form:

Powder and solvent for solution for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Cattle (calf)**
  - Meat. 9 day
- **Horse**
  - Meat. 16 day
- **Pig**
  - Meat. 9 day
- **Cattle**
  - Meat. 9 day
  - Milk. 3 day

Intravenous use:

- **Cattle**
  - Meat. 5 day
  - Milk. 24 hour
- **Cattle (calf)**
  - Meat. 5 day
- **Horse**
  - Meat. 5 day

Subcutaneous use:

- **Cattle**
  - Meat. 9 day
  - Milk. 3 day
- **Cattle (calf)**
  - Meat. 9 day
- **Pig**
  - Meat. 9 day
- **Horse**
  - Meat. 16 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01CA04

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

Authorisation status:
Valid

Authorised in:
Lithuania

Package description:
Available only in Lithuanian

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:
Bela-Pharm GmbH & Co. KG

Marketing authorisation date:
15/07/2007

Manufacturing sites for batch release:
Bela-Pharm GmbH & Co. KG

Responsible authority:
State Food And Veterinary Service

Authorisation number:
LT/2/07/1761/001

Date of authorisation status change:
13/07/2012

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.