AMOXYCILLIN 15% LA INJ.

Authorised

- Amoxicillin

Product identification

Medicine name:
AMOXYCILLIN 15% LA INJ.

Active substance:
Amoxicillin

Target species:
Cattle
Sheep
Pig
Dog
Cat

Route of administration:
Intramuscular use
Subcutaneous use

Product details

Active substance and strength:
Amoxicillin
150.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Suspension for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Cattle**
  - Meat and offal. 24 day
  - Milk. 3 day
- **Sheep**
  - Meat and offal. 16 day

Мляко: не се разрешава за употреба при животни, чието мляко е предназначено за консумация от хора

- **Pig**
  - Meat and offal. 16 day

Subcutaneous use:

- **Dog**
- **Cat**

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01CA04

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

Authorisation status:
Valid

Authorised in:
Bulgaria

Package description:
Available only in Bulgarian

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Well-established use application (Article 13a of Directive No 2001/82/EC)

Marketing authorisation holder:
Vazrajdane-Kasis OOD

Marketing authorisation date:
10/10/2019

Manufacturing sites for batch release:
Kepro B.V.

Responsible authority:
Bulgarian Food Safety Authority

Authorisation number:
0022-2931

Date of authorisation status change:
16/03/2023

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.
