Betamox LA Injection

Authorised

- Amoxicillin

Product identification

Medicine name:
Betamox LA Injection

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. 39 day
  - Milk. 108 hour Мляко: 108 часа (4.5 дни)
- **Sheep**
  - Meat and offal. 29 day

Не се разрешава за употреба при животни, чието мляко е предназначено за консумация от хора

- **Pig**
  - Meat and offal. 42 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01CA04

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

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Authorisation status:
Valid

Authorised in:
Bulgaria

Available in:
Bulgaria

Package description:
Available only in Bulgarian

Available only in Bulgarian

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Asklep-Pharma OOD

Marketing authorisation date:
10/10/2006

Manufacturing sites for batch release:
Norbrook Laboratories Limited
Norbrook Laboratories (Ireland) Limited

Responsible authority:
Bulgarian Food Safety Authority

Authorisation number:
0022-1658

Date of authorisation status change:
16/11/2011

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

Documents

Summary of Product Characteristics

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Package Leaflet and Labelling

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