AMOXYVET 500 mg/g POWDER FOR ORAL SOLUTION

Authorised

- Amoxicillin

Product identification

Medicine name:
АМОКСИВЕТ 500 mg/g ПРАХ ЗА ПЕРОРАЛЕН РАЗТВОР
AMOXYVET 500 mg/g POWDER FOR ORAL SOLUTION

Active substance:
Amoxicillin

Target species:
Cattle
Pig
Chicken (chick)

Route of administration:
In drinking water use

Product details

Active substance and strength:
Amoxicillin
500.00 milligram(s) / 1.00 gram(s)

Pharmaceutical form:
Powder for use in drinking water

Withdrawal period by route of administration:

In drinking water use:

- **Cattle**
  - Meat and offal. 1 day
- **Pig**
  - Meat and offal. 7 day
- **Chicken (chick)**
  - Meat and offal. 3 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

Package description:

Available only in Bulgarian
Available only in Bulgarian
Available only in Bulgarian
Available only in Bulgarian
Available only in Bulgarian
Available only in Bulgarian
Available only in Bulgarian
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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:
Biovet AD

Marketing authorisation date:
1/09/2009

Manufacturing sites for batch release:
Biovet AD

Responsible authority:
Bulgarian Food Safety Authority

Authorisation number:
0022-2387

Date of authorisation status change:
6/03/2023

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.

Package Leaflet and Labelling

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