AMOXYVET 500 mg/g POWDER FOR ORAL SOLUTION

Authorised

- Amoxicillin

Product identification

Medicine name:
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

Authorisation status:
Valid

Authorised in:
Bulgaria

Package description:
Available only in Bulgarian
Available only in Bulgarian
Available only in Bulgarian
Available only in Bulgarian
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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

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Package Leaflet and Labelling

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