Amoxy Active 697 mg/g Oral Powder for Pigs and Chickens

Authorised

- Amoxicillin trihydrate

Product identification

Medicine name:
Amoxy Active 697 mg/g Oral Powder for Pigs and Chickens

Active substance:
Amoxicillin trihydrate

Target species:
Chicken
Pig

Route of administration:
In drinking water use

Product details

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

Pharmaceutical form:
Oral powder

Withdrawal period by route of administration:

In drinking water use:

- **Chicken**
  - Meat and offal. 1 day
- **Pig**
  - Meat and offal. 2 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:

United Kingdom (Northern Ireland)

Available in:

United Kingdom (Northern Ireland)

Package description:

500 g Securitainer: white polypropylene container , covered with a low-density polyethylene lid.
5 kg Bucket: white polypropylene bucket provided with a polypropylene lid.
250 g Securitainer: white polypropylene container , covered with a low-density polyethylene lid.
2,5 kg Bucket: white polypropylene bucket provided with a polypropylene lid.
100 g Securitainer: white polypropylene container , covered with a low-density polyethylene lid.
1 kg Securitainer: white polypropylene container , covered with a low-density polyethylene lid.
1 kg Bucket: white polypropylene bucket provided with a polypropylene lid.

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:
Dopharma Research B.V.

Marketing authorisation date:
27/06/2014

Manufacturing sites for batch release:
Dopharma B.V.

Responsible authority:
The Veterinary Medicines Directorate

Authorisation number:
Vm 28365/3001

Date of authorisation status change:
1/12/2024

Reference member state:
Netherlands

Procedure number:
NL/V/0179/001

Concerned member states:
Austria Belgium Bulgaria Croatia Czechia Denmark Estonia Finland France Germany Greece Hungary Ireland Italy Latvia Lithuania Norway Poland Portugal Romania Slovakia Sweden United Kingdom (Northern Ireland)

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

Documents

Combined File of all Documents