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Octacillin, 800 mg/g powder for oral solution for chickens

Authorised

- Amoxicillin trihydrate

Product identification

Medicine name:
Octacillin, 800 mg/g powder for oral solution for chickens

Active substance:
Amoxicillin trihydrate

Target species:
Chicken

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:
Powder for use in drinking water

Withdrawal period by route of administration:

In drinking water use:

- **Chicken**

  - Meat and offal. 1 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:

Germany

Available in:

Germany

Package description:

Sachet 500 g. Sachets consist of a polyester layer on the outside, inside layers of aluminium and polyamide and an inner layer of polyethylene.
Sachet 250 g. Sachets consist of a white layer on the outside, inside different transparent layers, a sub-layer of aluminium and an inner layer of polyethylene.
Sachet 500 g. Sachets consist of a white layer on the outside, inside different transparent layers, a sub-layer of aluminium and an inner layer of polyethylene.
Sachet 250 g. Sachets consist of a polyester layer on the outside, inside layers of aluminium and polyamide and an inner layer of polyethylene.
Sachet 100 g. Sachets consist of a white layer on the outside, inside different transparent layers, a sub-layer of aluminium and an inner layer of polyethylene.
Sachet 100 g. Sachets consist of a polyester layer on the outside, inside layers of aluminium and polyamide and an inner layer of polyethylene.
Sachet 1.0 kg. Sachets consist of a white layer on the outside, inside different transparent layers, a sub-layer of aluminium and an inner layer of polyethylene.
Sachet 1.0 kg. Sachets consist of a polyester layer on the outside, inside layers of aluminium and polyamide and an inner layer of polyethylene.

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:
Eurovet Animal Health B.V.

Marketing authorisation date:
21/04/2010

Manufacturing sites for batch release:
Eurovet Animal Health B.V.

Responsible authority:
Federal Office Of Consumer Protection And Food Safety

Authorisation number:
401355.00.00

Date of authorisation status change:
31/10/2014

Reference member state:
Netherlands

Procedure number:
NL/V/0144/001

Concerned member states:
France Germany

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

Documents

Combined File of all Documents

English (PDF)
Published on: 13/06/2022
Updated on: 14/03/2026
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octacilin par.pdf