

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

Octacillin 800 mg/g poeder voor gebruik in het drinkwater voor kippen

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

Authorisation status:

Valid

Authorised in:

Netherlands

Available in:

Netherlands

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:

15/10/2009

Manufacturing sites for batch release:

Eurovet Animal Health B.V.

Responsible authority:

Medicines Evaluation Board

Authorisation number:

REG NL 104691

Date of authorisation status change:

29/01/2022

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)

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