

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

Authorisation status:

Valid

Authorised in:

France

Available in:

France

Package description:

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.

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 100 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 250 g. Sachets consist of a polyester layer on the outside, inside layers of aluminium and polyamide 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 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 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:

8/03/2010

Manufacturing sites for batch release:

Eurovet Animal Health B.V.

Responsible authority:

French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:

FR/V/0498260 5/2010

Date of authorisation status change:

8/03/2015

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

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.

octacilin par.pdf