Octacillin 800 mg/g powder for use in drinking water for pigs, amoxicillin trihydrate

Authorised

- Amoxicillin trihydrate

Product identification

Medicine name:
Octacillin 800 mg/g powder for use in drinking water for pigs, amoxicillin trihydrate
Octacillin 800 mg/g, milteliai naudoti su geriamuoju vandeniu kiaulėms

Active substance:
Amoxicillin trihydrate

Target species:
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:
Powder for use in drinking water

Withdrawal period by route of administration:

In drinking water use:

- **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:

Lithuania

Package description:

1 kg multilayer sachet consisting of the following materials: on the outside a white layer, inside different transparent layers, a sub-layer of aluminium and an inner layer of polyethylene

100 g multilayer sachet consisting of the following materials: on the outside a polyester layer, inside layers of aluminium and polyamide and an inner layer of polyethylene

500 g multilayer sachet consisting of the following materials: on the outside a polyester layer, inside layers of aluminium and polyamide and an inner layer of polyethylene

250 g multilayer sachet consisting of the following materials: on the outside a polyester layer, inside layers of aluminium and polyamide and an inner layer of polyethylene

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of polyethylene
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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:
17/05/2021

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

Responsible authority:
State Food And Veterinary Service

Authorisation number:
LT/2/21/2664/001-008

Date of authorisation status change:
17/05/2021

Reference member state:
Netherlands

Procedure number:
NL/V/0367/001

Concerned member states:
Austria Belgium Croatia Czechia Denmark France Germany Greece

Hungary Ireland Italy Latvia Lithuania Luxembourg Poland Portugal Romania Slovakia Slovenia Spain United Kingdom (Northern Ireland)

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

Documents

RV2664.pdf
