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

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

Authorisation status:
Valid

Authorised in:
Luxembourg

Package description:
500 g 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
250 g 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 white layer, inside different transparent layers, a sub-layer of aluminium and an inner layer of polyethylene
1 kg 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
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
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
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

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:
Dechra Regulatory B.V.

Marketing authorisation date:
13/12/2010

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

Responsible authority:
Ministere De La Sante Division De La Pharmacie Et Des Medicaments

Authorisation number:
V 914/10/12/2082

Date of authorisation status change:
13/12/2010

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)

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