Huvacillin 800 mg/g Powder for Use in Drinking Water for Chickens and Pigs

Authorised

- Amoxicillin trihydrate

Product identification

Medicine name:
Huvacillin 800 mg/g Powder for Use in Drinking Water for Chickens and Pigs

Active substance:
Amoxicillin trihydrate

Target species:
Chicken
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:

- **Chicken**
  - Meat and offal. 1 day

- **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:
United Kingdom (Northern Ireland)

Package description:
Zipped bag of 1kg made of low density polyethylene/aluminium/polyethylene terephthalate.
Zipped bag of 500 g made of low density polyethylene/aluminium/polyethylene terephthalate.
thermo-sealed sachet of 100 g made of low density polyethylene/aluminium/polyethylene terephthalate
Jar of 100 g made of high density polyethylene closed with a seal made of low density polyethylene/ polyethylene terephthalate/aluminium and a screw cap made of polypropylene

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:
HuVepharma

Marketing authorisation date:
25/08/2022

Manufacturing sites for batch release:
Huvepharma S.A.

Responsible authority:
The Veterinary Medicines Directorate

Authorisation number:
Vm 30282/3000

Date of authorisation status change:
28/04/2024

Reference member state:
Netherlands

Procedure number:
NL/V/0365/001

Concerned member states:
Austria Belgium Bulgaria Croatia Cyprus Czechia Denmark Estonia France Germany Greece Hungary Ireland Italy Latvia Lithuania 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