# Biocillin 1000 mg/g Powder for use in drinking water for chickens, ducks and turkeys

Authorised

- Amoxicillin trihydrate

## Product identification

**Medicine name:**
Biocillin 1000 mg/g Powder for use in drinking water for chickens, ducks and turkeys

**Active substance:**
Amoxicillin trihydrate

**Target species:**
Turkey
Chicken
Duck

**Route of administration:**
In drinking water use

## Product details

**Active substance and strength:**
Amoxicillin trihydrate
1000.00 milligram(s) / 1.00 gram(s)

**Pharmaceutical form:**
Powder for use in drinking water

**Withdrawal period by route of administration:**
**In drinking water use:**

- **Turkey**
  - Egg. no withdrawal period

Not authorised for use in birds producing eggs for human consumption and within 3 weeks of the start of the laying period.

  - Meat and offal. 5 day

- **Chicken**
  - Egg. no withdrawal period

Not authorised for use in birds producing eggs for human consumption and within 3 weeks of the start of the laying period.

  - Meat and offal. 1 day

- **Duck**
  - Meat and offal. 9 day
  - Egg. no withdrawal period

Not authorised for use in birds producing eggs for human consumption and within 3 weeks of the start of the laying period.

**Anatomical therapeutic chemical veterinary (ATCvet) codes:**
QJ01CA04

**Legal status of supply:**
Veterinary medicinal product subject to veterinary prescription

**Authorisation status:**
Valid

**Authorised in:**
Greece

**Available in:**
Greece

**Package description:**
(ID1) 250 gram(s): 1 Container (paper; polyethylene; aluminium; low-density polyethylene) with 250 gram(s)
(ID2) 500 gram(s): 1 Container (paper; polyethylene; aluminium; low-density polyethylene) with 500 gram(s)
(ID3) 1 kilogram(s): 1 Container (paper; polyethylene; aluminium; low-density polyethylene) with 1 kilogram(s)
(ID4) 2.5 kilogram(s): 1 Bag (polyethylene; paper; polyethylene; aluminium; polyethylene) with 2.5 kilogram(s)
(ID5) 5 kilogram(s): 1 Bag (polyethylene; paper; polyethylene; aluminium; polyethylene) with 5 kilogram(s)

## 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:**
Bela-Pharm GmbH & Co. KG

**Marketing authorisation date:**
25/02/2018

**Manufacturing sites for batch release:**
Bela-Pharm GmbH & Co. KG

**Responsible authority:**
National Organization For Medicines

**Authorisation number:**

105062/08-11-2021/K-0213501

**Date of authorisation status change:**
7/11/2021

**Reference member state:**
Germany

**Procedure number:**
DE/V/0182/001

**Concerned member states:**
Greece Ireland Poland United Kingdom (Northern Ireland)

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

## Documents

Combined File of all Documents

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