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Citramox 1000 mg/g powder for use in drinking water for chickens, ducks, turkeys and pigs

Authorised

- Amoxicillin trihydrate

Product identification

Medicine name:
Citramox 1000 mg/g powder for use in drinking water for chickens, ducks, turkeys and pigs

Active substance:
Amoxicillin trihydrate

Target species:
Turkey
Chicken
Duck
Pig

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**
  - Eggs. no withdrawal period

Do not use in birds in lay and within 3 weeks before the start of the laying period.

  - Meat and offal. 5 day

- **Chicken**
  - Eggs. no withdrawal period

Do not use in birds in lay and within 3 weeks before the start of the laying period.

  - Meat and offal. 1 day

- **Duck**
  - Eggs. no withdrawal period

Do not use in birds in lay and within 3 weeks before the start of the laying period.

  - Meat and offal. 9 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:
Cyprus

Package description:
Thermosealed bags made of polyester, aluminium and polyethylene complex.Pack sizes: 20 x 200 g
Thermosealed bags made of polyester, aluminium and polyethylene complex.Pack sizes: 1 kg bag
Thermosealed bags made of polyester, aluminium and polyethylene complex.Pack sizes: 500 g bag
Thermosealed bags made of polyester, aluminium and polyethylene complex.Pack sizes: 200 g bag

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Hybrid application (Article 13(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Laboratorios Karizoo S.A.

Marketing authorisation date:
11/02/2020

Manufacturing sites for batch release:
Laboratorios Karizoo S.A.

Responsible authority:
Veterinary Services, Ministry Of Agriculture, Natural Resources And Environment

Authorisation number:
CY00777V

Date of authorisation status change:
1/12/2021

Reference member state:
Ireland

Procedure number:
IE/V/0353/001

Concerned member states:
Cyprus Czechia Germany Greece Hungary Italy Latvia Lithuania Netherlands 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

Summary of Product Characteristics

Combined File of all Documents

English (PDF)
Published on: 13/07/2025
Updated on: 13/03/2026
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