Rhemox 1000, 1000 mg/g Powder for Use in Drinking Water for Chickens, Ducks, Turkeys

Authorised

- Amoxicillin trihydrate

Product identification

Medicine name:
Rhemox 1000, 1000 mg/g Powder for Use in Drinking Water for Chickens, Ducks, Turkeys
Rhemox Forte, 1000 mg/g poeder voor gebruik in drinkwater voor kippen, eenden, kalkoenen

Active substance:
Amoxicillin trihydrate

Target species:
Turkey
Duck
Chicken

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**
  - Meat and offal. 5 day
  - Eggs. no withdrawal period

Eggs: Not authorised for use in laying birds producing eggs for human consumption. Do not use within 3 weeks of onset of the laying period.

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

Eggs: Not authorised for use in laying birds producing eggs for human consumption. Do not use within 3 weeks of onset of the laying period.

- **Chicken**
  - Meat and offal. 1 day
  - Eggs. no withdrawal period

Eggs: Not authorised for use in laying birds producing eggs for human consumption. Do not use within 3 weeks of onset 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:
Netherlands

Package description:
sachet of 5 kg
sachet of 1 kg
sachet of 500 g
sachet of 200 g
sachet of 100 g

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:
Industrial Veterinaria S.A.

Marketing authorisation date:
22/06/2016

Manufacturing sites for batch release:
aniMedica GmbH
Industria Italiana Integratori Trei S.p.A.

Responsible authority:
Medicines Evaluation Board

Authorisation number:
REG NL 116268

Date of authorisation status change:
2/02/2022

Reference member state:
Spain

Procedure number:
ES/V/0331/001

Concerned member states:
Austria Belgium Germany Greece Hungary Italy Netherlands Poland Portugal Romania 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

This document does not exist in this language (English). You can find it in another language below.

Summary of Product Characteristics

Package Leaflet

eu-PUAR-rhemox-1000--1000-mg-g-powder-for-use-in-drinking-water-for-chickens--ducks--turkeys-en.pdf

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