

# Rhemox 500 mg/g powder for use in drinking water for pigs, chicken broilers, duck broilers and turkeys for meat production

Authorised

- Amoxicillin trihydrate

## Product identification

### **Medicine name:**

Rhemox 500 mg/g powder for use in drinking water for pigs, chicken broilers, duck broilers and turkeys for meat production

### **Active substance:**

Amoxicillin trihydrate

### **Target species:**

Pig

Chicken (broiler)

Turkey (for meat production)

Duck (broiler)

### **Route of administration:**

In drinking water use

## Product details

### **Active substance and strength:**

Amoxicillin trihydrate

500.00 milligram(s) / 1.00 gram(s)

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### **Pharmaceutical form:**

Powder for use in drinking water

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### **Withdrawal period by route of administration:**

#### **In drinking water use:**

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#### **Pig**

- Meat and offal. 6 day

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#### **Chicken (broiler)**

- Meat and offal. 1 day

- Eggs. no withdrawal period

Eggs: Not for use in birds producing or intended to produce eggs for human consumption. Do not use within 4 weeks of the start of the laying period.

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#### **Turkey (for meat production)**

- Meat and offal. 5 day

- Eggs. no withdrawal period

Eggs: Not for use in birds producing or intended to produce eggs for human consumption. Do not use within 4 weeks of the start of the laying period.

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#### **Duck (broiler)**

- Meat and offal. 9 day

- Eggs. no withdrawal period

Eggs: Not for use in birds producing or intended to produce eggs for human consumption. Do not use within 4 weeks of the start of the laying period.

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

QJ01CA04

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Greece

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**Available in:**

Greece

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**Package description:**

sachet of 1 kg

sachet of 400 g

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Generic application (Article 13(1) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Industrial Veterinaria S.A.

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**Marketing authorisation date:**

7/03/2016

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**Manufacturing sites for batch release:**

aniMedica Herstellungs GmbH  
Industria Italiana Integratori Trei S.p.A.  
Industrial Veterinaria S.A.

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**Responsible authority:**

National Organization For Medicines

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**Authorisation number:**

103208/21-10-2020/K-0178002

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**Date of authorisation status change:**

28/09/2023

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**Reference member state:**

Spain

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**Procedure number:**

ES/V/0236/001

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**Concerned member states:**

Austria Belgium Bulgaria Cyprus Czechia France Germany Greece Hungary  
Ireland Italy Poland Portugal Romania Slovakia Slovenia  
United Kingdom (Northern Ireland)

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To consult adverse reactions on veterinary medicinal products please go to  
[www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Summary of Product Characteristics

English (PDF)

Published on: 5/04/2023

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Package Leaflet

English (PDF)

Published on: 5/04/2023

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Labelling

Combined File of all Documents

eu-PUAR-esv0236001-mrp-rhemox-500-mg-g-en.pdf

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