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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)

Pharmaceutical form:
Powder for use in drinking water

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

- **Pig**
  - Meat and offal. 6 day

- **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.

- **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.

- **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.

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01CA04

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

Authorisation status:
Valid

Authorised in:
Bulgaria

Available in:
Bulgaria

Package description:
sachet of 400 g
sachet of 1 kg

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:
10/09/2015

Manufacturing sites for batch release:

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

Responsible authority:
Bulgarian Food Safety Authority

Authorisation number:
0022-2606

Date of authorisation status change:
10/09/2015

Reference member state:
Spain

Procedure number:
ES/V/0236/001

Concerned member states:
Austria Belgium Bulgaria Cyprus Czechia France Germany Greece Hungary Ireland Italy Poland Portugal Romania Slovakia Slovenia

United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics

English (PDF)
Published on: 14/03/2026
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Package Leaflet

English (PDF)
Published on: 22/06/2026
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Labelling

English (PDF)
Published on: 27/06/2024
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Combined File of all Documents

Package Leaflet and Labelling

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

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