

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

Authorised

- Amoxicillin trihydrate

Product identification

Medicine name:

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

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:

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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.

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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.

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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:

United Kingdom (Northern Ireland)

Available in:

United Kingdom (Northern Ireland)

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:

14/06/2016

Manufacturing sites for batch release:

aniMedica GmbH

Industria Italiana Integratori Trei S.p.A.

Responsible authority:

The Veterinary Medicines Directorate

Authorisation number:

Vm 36547/3002

Date of authorisation status change:

20/10/2024

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

Summary of Product Characteristics

Package Leaflet

Combined File of all Documents

Labelling

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