SURAMOX 1000 MG/G POWDER FOR USE IN DRINKING WATER FOR CHICKENS, DUCKS, TURKEYS

Authorised

- Amoxicillin trihydrate

Product identification

Medicine name:
SURAMOX 1000 MG/G POWDER FOR USE IN DRINKING WATER FOR CHICKENS, DUCKS, TURKEYS

Active substance:
Amoxicillin trihydrate

Target species:
Turkey
Duck
Chicken
Chicken (broiler)

Route of administration:
Oral 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:
Oral use:

- **Turkey**
  - Eggs. no withdrawal period

The product is not authorised for use in laying birds producing eggs for human consumption and within 3 weeks of onset of laying.

  - Meat and offal. 5 day

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

The product is not authorised for use in laying birds producing eggs for human consumption and within 3 weeks of onset of laying.

  - Meat and offal. 9 day

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

The product is not authorised for use in laying birds producing eggs for human consumption and within 3 weeks of onset of laying.

- **Chicken (broiler)**
  - Meat and offal. 1 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01CA04

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Germany

Available in:
Germany

Package description:
Available only in French
Available only in French
Available only in French
Available only in French

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:
Virbac

Marketing authorisation date:
2/09/2013

Manufacturing sites for batch release:
aniMedica Herstellungs GmbH

Responsible authority:
Federal Office Of Consumer Protection And Food Safety

Authorisation number:
401891.00.00

Date of authorisation status change:
16/10/2018

Reference member state:
France

Procedure number:
FR/V/0370/001

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

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

Combined File of all Documents