SURRICOXX 400 MG/ML SOLUTION FOR USE IN DRINKING WATER FOR CHICKENS, TURKEYS, DUCKS, AND GUINEA FOWLS

Authorised

- Amprolium hydrochloride

Product identification

Medicine name:
SURRICOXX 400 MG/ML SOLUTION FOR USE IN DRINKING WATER FOR CHICKENS, TURKEYS, DUCKS, AND GUINEA FOWLS
SURRICOXX 400 mg/ml разтвор за прилагане във водата за пиене за пилета, пуйки, патици и токачки

Active substance:
Amprolium hydrochloride

Target species:
Turkey
Guinea fowl
Duck
Chicken

Route of administration:
In drinking water use

Product details

Active substance and strength:
Amprolium hydrochloride
452.40 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for use in drinking water

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

- **Turkey**
  - Meat and offal. 0 day
  - Eggs. 0 day
- **Guinea fowl**
  - Meat and offal. 0 day
  - Eggs. 0 day
- **Duck**
  - Meat and offal. 0 day
  - Eggs. 0 day
- **Chicken**
  - Meat and offal. 0 day
  - Eggs. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QP51AX09

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Bulgaria

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

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Hybrid application (Article 13(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
V.M.D.

Marketing authorisation date:
24/01/2021

Manufacturing sites for batch release:
LABORATOIRES BIOVE
VMD N.V.

Responsible authority:
Bulgarian Food Safety Authority

Authorisation number:
0022-3036

Date of authorisation status change:
24/01/2021

Reference member state:
France

Procedure number:
FR/V/0422/001

Concerned member states:
Austria Belgium Bulgaria Czechia Estonia Germany Greece Hungary Ireland Italy Latvia Lithuania Luxembourg Malta Netherlands Poland Portugal Romania Slovakia 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

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Package Leaflet and Labelling

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