AMPROLINE 400 MG/ML SOLUTION FOR USE IN DRINKING WATER FOR CHICKENS AND TURKEYS

Authorised

- Amprolium hydrochloride

Product identification

Medicine name:
AMPROLINE 400 MG/ML SOLUTION FOR USE IN DRINKING WATER FOR CHICKENS AND TURKEYS
Amproline 400 mg/ml Oplossing voor gebruik in drinkwater
Amproline 400 mg/ml Solution pour administration dans l'eau de boisson
Amproline 400 mg/ml Lösung zum Eingeben über das Trinkwasser

Active substance:
Amprolium hydrochloride

Target species:
Turkey
Chicken (pullet for egg production, future layer)
Chicken (for reproduction)
Chicken (broiler)
Chicken (layer hen)

Route of administration:
Oral 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:
Oral use:

- **Turkey**
  - Eggs. 0 day
  - Meat and offal. 0 day
- **Chicken (pullet for egg production, future layer)**
  - Eggs. 0 day
  - Meat and offal. 0 day
- **Chicken (for reproduction)**
  - Eggs. 0 day
  - Meat and offal. 0 day
- **Chicken (broiler)**
  - Meat and offal. 0 day
- **Chicken (layer hen)**
  - Eggs. 0 day
  - Meat and offal. 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:
Belgium

Available in:
Belgium

Package description:
1 L can
5 L can
100 mL can

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:
Huvepharma S.A.

Marketing authorisation date:
16/09/2019

Manufacturing sites for batch release:
Huvepharma S.A.
Biovet AD

Responsible authority:

Federal Agency For Medicines And Health Products

Authorisation number:
This information is not available for this product.

Date of authorisation status change:
16/09/2019

Reference member state:
France

Procedure number:
FR/V/0284/001

Concerned member states:
Austria Belgium Bulgaria Czechia Denmark Germany Greece Hungary Ireland Italy Netherlands Poland Portugal Romania 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.

Package Leaflet

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

Labelling

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