File downloaded on 2026-11-04
Source URL: https://medicines.health.europa.eu/veterinary/en/600000043593

COCCIBAL 200 MG/ML SOLUTION FOR USE IN DRINKING WATER FOR CHICKENS AND TURKEYS

Authorised

- Amprolium hydrochloride

Product identification

Medicine name:
COCCIBAL 200 MG/ML SOLUTION FOR USE IN DRINKING WATER FOR CHICKENS AND TURKEYS

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
226.20 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for use in drinking water

Withdrawal period by route of administration:
Oral use:

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QP51BX02

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

Authorisation status:
Valid

Authorised in:
Denmark

Available in:
Denmark

Package description:
1 L bottle
Box of 10 bottles of 100 mL
Box of 4 barrels of 5 L
Box of 12 bottles of 1 L
5 L barrel

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:
S P Veterinaria S.A.

Marketing authorisation date:
25/04/2012

Manufacturing sites for batch release:
S P Veterinaria S.A.

Responsible authority:
Danish Medicines Agency

Authorisation number:
47846

Date of authorisation status change:
25/04/2012

Reference member state:
France

Procedure number:
FR/V/0230/001

Concerned member states:
Belgium Bulgaria Cyprus Czechia Denmark Germany Greece Hungary Ireland Italy Luxembourg Malta 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

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

Combined File of all Documents

eu-puar-frv0230001-mr-rpe291-en.pdf