VETMULIN 125 mg/ml Solution for use in drinking water for pigs and chickens

Authorised

- Tiamulin hydrogen fumarate
- Tiamulin hydrogen fumarate

Product identification

Medicine name:
VETMULIN 125 mg/ml Solution for use in drinking water for pigs and chickens

Active substance:
Tiamulin hydrogen fumarate

Tiamulin hydrogen fumarate

Target species:
Poultry
Pig

Route of administration:
In drinking water use

Product details

Active substance and strength:
Tiamulin hydrogen fumarate

125.00 milligram(s) / 1.00 millilitre(s)

Tiamulin hydrogen fumarate
125.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for use in drinking water

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

- **Poultry**
  - Meat and offal. 2 day
  - Meat and offal. 2 day
  - Eggs. 0 day
  - Eggs. 0 day
- **Pig**
  - Meat and offal. 2 day

8,8 mg tiamulinhydrogenfumarat/kg kropsvægt svarende til 7 ml af produktet/100 kg kropsvægt

- Meat and offal. 2 day

8,8 mg tiamulinhydrogenfumarat/kg kropsvægt svarende til 7 ml af produktet/100 kg kropsvægt

- Meat and offal. 4 day

20 mg tiamulinhydrogenfumarat/kg kropsvægt svarende til 16 ml af produktet/100 kg kropsvægt

- Meat and offal. 4 day

20 mg tiamulinhydrogenfumarat/kg kropsvægt svarende til 16 ml af produktet/100 kg kropsvægt

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Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01XQ01

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

Authorisation status:
Valid

Authorised in:
Greece

Package description:
Available only in Danish
Available only in Danish

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:
HuVepharma

Marketing authorisation date:
27/02/2019

Manufacturing sites for batch release:
Biovet AD

Responsible authority:
National Organization For Medicines

Authorisation number:
21524/28-02-2019/K-0170507

Date of authorisation status change:
24/11/2020

Reference member state:

Denmark

Procedure number:
DK/V/0122/001

Concerned member states:
Austria Belgium Bulgaria Croatia Cyprus Czechia Estonia France Germany Greece Hungary Ireland Italy Latvia Lithuania Netherlands Poland Portugal Romania Slovakia Slovenia 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.