

# 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

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**Active substance:**

Tiamulin hydrogen fumarate

Tiamulin hydrogen fumarate

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**Target species:**

Poultry

Pig

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**Route of administration:**

In drinking water use

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## 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)

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**Pharmaceutical form:**

Solution for use in drinking water

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**Withdrawal period by route of administration:**

**In drinking water use:**

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**Poultry**

- Meat and offal. 2 day
- Meat and offal. 2 day
- Eggs. 0 day
- Eggs. 0 day

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**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

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Slovenia

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**Available in:**

Slovenia

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**Package description:**

Available only in [Danish](#)

Available only in [Danish](#)

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Generic application (Article 13(1) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

HuVepharma

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**Marketing authorisation date:**

4/02/2019

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**Manufacturing sites for batch release:**

Biovet AD

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**Responsible authority:**

Agency For Medicinal Products And Medical Devices Of The Republic Of Slovenia

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**Authorisation number:**

DC/V/0651/001

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**Date of authorisation status change:**

4/02/2019

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**Reference member state:**

Denmark

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**Procedure number:**

DK/V/0122/001

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**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)

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To consult adverse reactions on veterinary medicinal products please go to  
[www.adrreports.eu/vet](http://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.

PI.pdf