

# Doxylin 100%, powder for use in drinking water/milk for calves and pigs

Authorised

- Doxycycline hyclate

## Product identification

**Medicine name:**

Doxylin 100%, powder for use in drinking water/milk for calves and pigs

---

**Active substance:**

Doxycycline hyclate

---

**Target species:**

Cattle (calf)

Pig

---

**Route of administration:**

In drinking water/milk use

---

## Product details

**Active substance and strength:**

Doxycycline hyclate

1000.00 milligram(s) / 1.00 gram(s)

---

**Pharmaceutical form:**

Powder for use in drinking water/milk

---

**Withdrawal period by route of administration:****In drinking water/milk use:**

- 

**Cattle (calf)**

- Meat and offal. 14 day

- 

**Pig**

- Meat and offal. 8 day

---

**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QJ01AA02

---

**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

---

**Authorisation status:**

Valid

---

**Authorised in:**

Lithuania

---

**Package description:**

White polypropylene container (bucket) of 5 kg, covered with a polypropylene closure.

White polypropylene container (bucket) of 2 kg, covered with a polypropylene closure.

White polypropylene container of 100 grams, covered with a low-density polyethylene closure.

White polypropylene container (bucket) of 1 kg, covered with a polypropylene closure.

White polypropylene container of 1 kg, covered with a low-density polyethylene closure.

---

## 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:**

Dopharma Research B.V.

---

**Marketing authorisation date:**

17/12/2013

---

**Manufacturing sites for batch release:**

Dopharma B.V.

---

**Responsible authority:**

State Food And Veterinary Service

---

**Authorisation number:**

LT/2/13/2203/001-005

---

**Date of authorisation status change:**

28/01/2026

---

**Reference member state:**

Netherlands

---

**Procedure number:**

NL/V/0184/001

---

**Concerned member states:**

Belgium Denmark Estonia France Germany Greece Hungary Italy Latvia  
Lithuania Poland Romania

---

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

## Documents

Combined File of all Documents

English (PDF)

Published on: 28/01/2026

[Download](#)