

File downloaded on 2026-06-11

**Source URL:** <https://medicines.health.europa.eu/veterinary/en/600000014072>

# LINCOMICINA FP 110 mg/g

Authorised

- Lincomycin

## Product identification

**Medicine name:**

LINCOMICINA FP 110 mg/g

---

**Active substance:**

Lincomycin

---

**Target species:**

Pig  
Chicken (hen)  
Turkey  
Duck  
Goose  
Pigeon

---

**Route of administration:**

In-feed use

---

## Product details

**Active substance and strength:**

Lincomycin  
11.00 gram(s) / 100.00 gram(s)

---

**Pharmaceutical form:**

Premix for medicated feeding stuff

---

**Withdrawal period by route of administration:****In-feed use:**

- 

**Pig**

- Meat and offal. 4 day

- 

**Chicken (hen)**

- Meat and offal. 5 day

- Eggs. 6 day

- 

**Turkey**

- Meat and offal. 7 day

Nu se administrează la curcile ale căror ouă sunt destinate consumului uman.

- 

**Duck**

- Meat and offal. 5 day

Nu se administrează la rațele ale căror ouă sunt destinate consumului uman.

- 

**Goose**

- Meat and offal. 5 day

Nu se administrează la găștele ale căror ouă sunt destinate consumului uman.

- 

**Pigeon**

- Meat and offal. 5 day

---

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

QJ01FF02

---

**Legal status of supply:**

This information is not available for this product.

---

**Authorisation status:**

Valid

---

**Authorised in:**

Romania

---

**Available in:**

Romania

---

**Package description:**

Available only in Romanian

Available only in Romanian

Available only in Romanian

Available only in Romanian

Available only in Romanian

Available only in Romanian

Available only in Romanian

Available only in Romanian

Available only in Romanian

Available only in Romanian

Available only in Romanian

Available only in Romanian

Available only in Romanian

Available only in Romanian

Available only in Romanian

Available only in Romanian

Available only in Romanian

Available only in Romanian

---

## Additional information

**Entitlement type:**

Marketing Authorisation

---

**Legal basis of product authorisation:**

Complete application (stand-alone) - Directive No 2001/82/EC

---

**Marketing authorisation holder:**

Pasteur Filiala Filipesti S.A.

---

**Marketing authorisation date:**

10/10/2000

---

**Manufacturing sites for batch release:**

Pasteur Filiala Filipesti S.A.

---

**Responsible authority:**

Institute For Control Of Biological Products And Veterinary Medicines

---

**Authorisation number:**

130135

---

**Date of authorisation status change:**

27/08/2025

---

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

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