

# Gallifen 40 mg/g premix for medicated feeding stuff for chickens and pheasants

Authorised

- Fenbendazole

## Product identification

**Medicine name:**

Gallifen 40 mg/g premix for medicated feeding stuff for chickens and pheasants

---

**Active substance:**

Fenbendazole

---

**Target species:**

Chicken

Pheasant

---

**Route of administration:**

In-feed use

---

## Product details

**Active substance and strength:**

Fenbendazole

40.00 milligram(s) / 1.00 gram(s)

---

**Pharmaceutical form:**

Premix for medicated feeding stuff

---

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

- 

**Chicken**

- Eggs. 0 day
- Meat and offal. 8 day

- 

**Pheasant**

- Eggs. 0 day
- Meat and offal. 8 day

Do not release pheasants for hunting for at least 8 days after the end of medication

---

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

QP52AC13

---

**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

---

**Authorisation status:**

Valid

---

**Authorised in:**

Belgium

---

**Available in:**

Belgium

---

**Package description:**

Polyethylene/aluminium foil/polyethylene terephthalate zipper bags of 5 kg.  
Polyethylene/aluminium foil/polyethylene terephthalate zipper bags of 2 kg.  
Polyethylene/aluminium foil/polyethylene terephthalate zipper bags of 1 kg.  
Polyethylene-aluminium-paper /paper/paper bag of 20 kg.

---

## 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:**

HuVepharma

---

**Marketing authorisation date:**

8/03/2017

---

**Manufacturing sites for batch release:**

Biovet AD

---

**Responsible authority:**

Federal Agency For Medicines And Health Products

---

**Authorisation number:**

This information is not available for this product.

---

**Date of authorisation status change:**

9/06/2022

---

**Reference member state:**

Ireland

---

**Procedure number:**

IE/V/0555/001

---

**Concerned member states:**

Belgium Bulgaria France Hungary Italy Netherlands Poland Portugal  
Romania Spain United Kingdom (Northern Ireland)

---

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

English (PDF)

Published on: 25/09/2024

[Download](#)

### 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.