

# 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  
ГАЛИФЕН 40 mg/g премикс за медикаментозен фураж за пилета и фазани

### **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

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

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**Chicken**

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

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**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

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

QP52AC13

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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:**

Bulgaria

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**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.

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Hybrid application (Article 13(3) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

HuVepharma

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**Marketing authorisation date:**

19/12/2018

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**Manufacturing sites for batch release:**

Biovet AD

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**Responsible authority:**

Bulgarian Food Safety Authority

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**Authorisation number:**

0022-2857

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**Date of authorisation status change:**

19/12/2018

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**Reference member state:**

Ireland

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**Procedure number:**

IE/V/0555/001

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**Concerned member states:**

Belgium Bulgaria France Hungary Italy Netherlands Poland Portugal  
Romania 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

English (PDF)

Published on: 25/09/2024

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### Package Leaflet

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### Labelling

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