

# Gallifen 200 mg/ml suspension for use in drinking water for chickens and pheasants

Authorised

- Fenbendazole

## Product identification

**Medicine name:**

Gallifen 200 mg/ml suspension for use in drinking water for chickens and pheasants

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**Active substance:**

Fenbendazole

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**Target species:**

Chicken

Pheasant

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**Route of administration:**

In drinking water use

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## Product details

**Active substance and strength:**

Fenbendazole

200.00 milligram(s) / 1.00 millilitre(s)

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**Pharmaceutical form:**

Suspension for use in drinking water

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

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**Chicken**

- Eggs. 0 day
- Meat and offal. 8 day \* when used at 3 mg fenbendazole / kg bw / day
- Meat and offal. 6 day

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**Pheasant**

- Eggs. 0 day
- Meat and offal. 6 day

Do not release pheasants for hunting for at least 6 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:**

Hungary

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**Available in:**

Hungary

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**Package description:**

White rectangular HDPE bottle of 1 litre with vertically see-through bar with an LDPE insert closed with white PP tamper-evident screw cap with a LDPE sealing disk.

White HDPE canisters with white HDPE ribbed tamper-evident screw cap of 5 litres.

White HDPE canisters with white HDPE ribbed tamper-evident screw cap of 2.5 litres.  
White cylindrical High Density Polyethylene (HDPE) bottle with white polypropylene (PP) screw tamper-evident closure of 1 litre.  
White cylindrical High Density Polyethylene (HDPE) bottle with white polypropylene (PP) screw tamper-evident closure of 125 ml.

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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:**

11/04/2018

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**Manufacturing sites for batch release:**

Biovet AD

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**Responsible authority:**

Directorate Of Veterinary Medicinal Products

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**Authorisation number:**

3954/X/18/NÉBIH ÁTI

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**Date of authorisation status change:**

11/04/2018

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**Reference member state:**

Ireland

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**Procedure number:**

IE/V/0579/001

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**Concerned member states:**

Austria Belgium Bulgaria Croatia Cyprus Czechia Denmark Estonia France  
Germany Greece Hungary Italy Latvia Lithuania Luxembourg 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