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
Gallifen 200 mg/ml Suspensie voor gebruik in drinkwater
Gallifen 200 mg/ml Suspension pour administration dans l'eau de boisson
Gallifen 200 mg/ml Suspension zum Eingeben über das Trinkwasser

Active substance:
Fenbendazole

Target species:
Chicken
Pheasant

Route of administration:
In drinking water use

Product details

Active substance and strength:
Fenbendazole
200.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Suspension for use in drinking water

Withdrawal period by route of administration:

In drinking water use:

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QP52AC13

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Belgium

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.

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:
29/03/2018

Manufacturing sites for batch release:
Biovet J.S.C.

Responsible authority:
Federal Agency For Medicines And Health Products

Authorisation number:
This information is not available for this product.

Date of authorisation status change:
26/10/2021

Reference member state:
Ireland

Procedure number:
IE/V/0579/001

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)

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Summary of Product Characteristics

English (PDF)
Published on: 26/05/2024
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Package Leaflet

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