Fluboral 200 mg/ml, suspension for use in drinking water for pigs and chickens

Authorised

- Flubendazole

Product identification

Medicine name:
Fluboral 200 mg/ml Suspension zum Eingeben über das Trinkwasser für Schweine und Hühner
Fluboral 200 mg/ml, suspension for use in drinking water for pigs and chickens

Active substance:
Flubendazole

Target species:
Chicken
Pig

Route of administration:
In drinking water use

Product details

Active substance and strength:
Flubendazole

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. 2 day
- **Pig**
  - Meat and offal. 4 day 1 mg / kg for 5 days.
  - Meat and offal. 5 day 2.5 mg / kg for 2 days.

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QP52AC12

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

Authorisation status:
Valid

Authorised in:
Germany

Package description:
Semi-transparent high density polyethylene bottle of 3 litre closed with white high density polyethylene screw-cap containing low density polyethylene sealing element.
Semi-transparent high density polyethylene bottle of 1 litre closed with white high density polyethylene screw-cap containing low density polyethylene sealing element.
Semi-transparent high density polyethylene bottle of 250 ml closed with white high density polyethylene screw-cap containing low density polyethylene sealing element.

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:
Dechra Regulatory B.V.

Marketing authorisation date:
13/01/2023

Manufacturing sites for batch release:
Genera d.d.

Responsible authority:
Federal Office Of Consumer Protection And Food Safety

Authorisation number:
V7007884.00.00

Date of authorisation status change:
13/01/2023

Reference member state:
Ireland

Procedure number:
IE/V/0664/001

Concerned member states:
Austria Belgium Croatia Denmark France Germany Italy Netherlands Poland Portugal Slovenia Spain United Kingdom (Northern Ireland)

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

Documents

Combined File of all Documents

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