FLIMABEND 100 MG/G SUSPENSION FOR USE IN DRINKING WATER FOR CHICKENS AND PIGS

Authorised

- Flubendazole

Product identification

Medicine name:
FLIMABEND 100 MG/G SUSPENSION FOR USE IN DRINKING WATER FOR CHICKENS AND PIGS

Active substance:
Flubendazole

Target species:
Chicken (broiler)
Pig
Chicken

Route of administration:
Oral use

Product details

Active substance and strength:
Flubendazole
100.00 milligram(s) / 1.00 gram(s)

Pharmaceutical form:
Oral suspension

Withdrawal period by route of administration:
Oral use:

- **Chicken (broiler)**
  - Meat and offal. 2 day
- **Pig**
  - Meat and offal. 3 day — Dose 1 mg/kg body weight for 5 days
  - Meat and offal. 4 day — Dose 2.5 mg/kg body weight for 2 days
- **Chicken**
  - Eggs. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QP52AC12

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

Authorisation status:
Valid

Authorised in:
Slovenia

Available in:
Slovenia

Package description:
Cardboard box containing 2 sachets of 20 g suspension for use in drinking water
Cardboard box containing 6 containers of 750 g suspension for use in drinking water
Cardboard box containing 4 containers of 750 g suspension for use in drinking water
Cardboard box containing 25 sachets of 100 g suspension for use in drinking water
Cardboard box containing 5 sachets of 100 g suspension for use in drinking water
Cardboard box containing 1 sachet of 100 g suspension for use in drinking water
Cardboard box containing 24 sachets of 50 g suspension for use in drinking water
Cardboard box containing 2 sachets of 50 g suspension for use in drinking water
Cardboard box containing 24 sachets of 20 g suspension for use in drinking water

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:
KRKA tovarna zdravil d.d. Novo mesto

Marketing authorisation date:
16/01/2013

Manufacturing sites for batch release:
TAD Pharma GmbH
KRKA tovarna zdravil d.d. Novo mesto

Responsible authority:
Agency For Medicinal Products And Medical Devices Of The Republic Of Slovenia

Authorisation number:
DC/V/0421/001

Date of authorisation status change:

16/01/2013

Reference member state:
France

Procedure number:
FR/V/0242/001

Concerned member states:
Austria Belgium Bulgaria Czechia Denmark Estonia Germany Greece Hungary Ireland Italy Latvia Lithuania Netherlands Norway Poland Portugal Romania Slovakia Slovenia Spain Sweden United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics

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.

Package Leaflet

This document does not exist in this language (English). You can find it in another language below.