

Consumer leaflet for a Veterinary Product

This medicine is marketed according to a veterinarian's prescription only.

FOR ANIMAL USE ONLY

1. Name of the veterinary medicine, form and strength

Flubengal 5% Veterinary
Powder for premix

2. Active ingredients

Flubendazole 5%

For the list of non-active materials, see section 13

3. What is the medicine intended for?

For the treatment of round and flat worms in the digestive system as *Ascaridia galli*, *Capillaria* spp, *Heterakis gallinarum*, *Raillietina* spp, in broilers and layers

Therapeutic group:

Anthelmintics Benzimidazoles group.

4. Contraindications

Unknown.

5. Side effects

Side effects can be reported to the Ministry of Health by clicking on the link "Adverse Drug Reactions Report" that appears on the home page of the Ministry of Health web site (www.health.gov.il), which leads to an online form for reporting side effects. Alternatively, you can use the following link: <https://sideeffects.health.gov.il/>

6. Target animals

broilers and layers.

7. Dosage and administration

600 gram flubengal (30 ppm flubendazole) per ton feed.

First mix with 5 kg feed and then with the rest of feed to make 1 ton. Mix the product thoroughly into the feed.

Administer the medicated feed for 7 consecutive days.

8. How to use the product

For administration in feed according to the instructions in section 7.

9. Withdrawal period

Slaughter for meat: 14 days

Eggs marketing: 1 day

10. Warnings

Avoid strategies that may lead to increased risk for development of resistance to anthelmintic medicines, which include:

- Frequent and repeated use of anthelmintic from the same class for a long period.
- Underdosing

Special warnings regarding the safety of the person administering the veterinary medicinal product

Contact with the product may cause sensitivity in the skin and eyes

People with known hypersensitivity (allergy) to flubendazole should avoid contact with the veterinary product

- When mixing the product in the feed, avoid contact of the medicine with skin and eyes and inhaling it, by using proper protective means like goggles, gloves and mask.
- Do not smoke, eat or drink when working with the product.

Wash hands after use of the product and after contact with the medicated feed.

In case of contact with skin or eyes, wash immediately with plenty of water.

If symptoms appear following exposure such as skin rash, consult the doctor and take the leaflet or label with you. Inflammation of the face, lips, eyes or breathing distress are severe symptoms which need urgent medical treatment.

11. Storage instructions

- Avoid poisoning! This medicine, and any other medicine, must be kept in a closed place outside the reach and sight of children and/or infants to prevent accidental poisoning.
- Do not use the medicine after its expiration date (exp. date) as it appears on the package. The expiration date refers to the last day of the stated month.
- Storage conditions: below 25°C.

12. Instructions for disposing the product / remaining product at the end of its use

Any residue of a veterinary medical product or any waste obtained from the use of a veterinary medical product should be disposed of as toxic waste, do not throw into the sewage system.

13. Additional Information

- In addition to the active ingredient, the product contains lactose
- What does the medicine look like and what is the content of the package – white to yellowish powder.
- Package sizes: 600 gram / 3 kg in laminate bag.

Manufacturer and License holder: BIOVAC LTD.,

Hailan Str., 30, OR-AKIVA 3060000

Registration number of this medicine in the Ministry of Health State Medicine Registry: 082-83-92321

Revised on 07.2023 in accordance with the guidelines of the Ministry of Health