

CANIVERM oral paste

1. NAME AND ADDRESS OF THE MARKETING AUTHORISATION HOLDER AND OF THE MANUFACTURING AUTHORISATION HOLDER RESPONSIBLE FOR BATCH RELEASE, IF DIFFERENT

Marketing authorisation holder and
manufacturer:

Bioveta a.s.
Komenského 212/12
683 23 Ivanovice na Hané
Czech Republic

2. NAME OF THE VETERINARY MEDICINAL PRODUCT

CANIVERM oral paste
Fenbendazole, pyrantel embonate,
praziquantel

3. STATEMENT OF THE ACTIVE SUBSTANCES AND OTHER SUBSTANCES

Yellow-brown paste with noticeable brown
particles of aroma contains in 1 ml:

Active substance(s):

Fenbendazole	75 mg
Pyrantel embonate	72 mg
Praziquantel	25 mg

Excipients:

Butylated hydroxyanisole	0.2 mg
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4. INDICATION(S)

Diseases caused by helminths in dogs and
cats (*Toxocara canis*, *Toxocara cati*, *Toxascaris
leonina*, *Uncinaria stenocephala*, *Ancylostoma
caninum*, *Trichuris vulpis*, *Echinococcus
granulosus*, *Echinococcus multilocularis*,
Dipylidium caninum, *Taenia spp.*, *Multiceps
multiceps*, *Mesocostoides spp.*).

5. CONTRAINDICATIONS

Ocular cysticercosis and spinal
neurocysticercosis. Liver impairment.
Do not administer concurrently with piperazine-
based products, or other antiparasitics.
Do not use in case of hypersensitivity to the
active substance or to any of the excipients.

6. ADVERSE REACTIONS

Hypersalivation, vomiting, mushy stools,
urticaria, or fever may occur within a few hours
after administration. These are reactions to
proteins released from dying helminths, not
reactions to the drug.

The product can cause drowsiness in animals.

If you notice any side effects, even those not
already listed in this package leaflet or you
think that the medicine has not worked, please
inform your veterinary surgeon.

7. TARGET SPECIES

Dogs and cats.

8. DOSAGE FOR EACH SPECIES, ROUTE(S) AND METHOD OF ADMINISTRATION

Dosage:

The recommended dose is 15 mg
fenbendazole, 14.4 mg pyrantel pamoate and
5 mg praziquantel per 1 kg bw, i.e. 1 ml of the
paste per 5 kg bw.

Cat:

- 0.5 ml of paste per 0.5-2 kg bw

- 1 ml of paste per 2.1-5 kg bw

Dog:

- 0.5 ml of paste per 0.5-2 kg bw

- 1 ml of paste per 2.1-5 kg bw

- then 1 ml of paste per every 5 kg bw

CANIVERM oral paste is intended for oral
administration only.

9. ADVICE ON CORRECT ADMINISTRATION

The applicator should be adjusted to the
calculated dose according to the scheme
above by setting the ring on a respective piston
position: Hold the piston of applicator, turn the
grooved dosing ring on the piston so that the
bottom edge of the ring is aligned with the mark
of the desired volume.

Make sure that the animal has no feed
residues in its mouth. Remove the cap from the
applicator, insert it into the animal's mouth and
apply the paste to the root of the tongue. After
application lift the animal's head immediately
for a few seconds and make sure that the
animal has swallowed the dose.

Do not administer with milk food.

Administer a single dose.

It is recommended to deworm young animals
once every 3 weeks from 3 to 12 weeks of age
and then regularly every 3 months.

10. WITHDRAWAL PERIOD

Not intended for food-producing animals.

11. SPECIAL STORAGE PRECAUTIONS

Keep out of the sight and reach of children.

This veterinary medicinal product does not
require any special storage conditions.

Do not use this veterinary medicinal product
after the expiry date which is stated on the
label and carton.

Shelf life after first opening the container:
6 months.

12. SPECIAL WARNING(S)

Special precautions for use in animals:

Dosing should be strictly observed in pregnant
bitches.

Possible side effects should be discussed with
a veterinarian.

The product must not be administered to
animals showing signs of other than intestinal
parasitic disease.

Parasite resistance against any group of
anthelmintics may develop following frequent,
repeated use of anthelmintics of the same
group.

Avoid underdosing due to incorrect
determination of body weight or improper
administration of the drug.

Special precautions to be taken by the person administering the veterinary medicinal product to animals:

The product may cause skin and eye irritation.
Avoid contact with skin and eyes. In case of
accidental contamination, wash the affected
area under a stream of clean water. In case of
accidental ingestion or eye contact seek
medical advice immediately and show the
package leaflet or the label to the physician.
Pregnant women and people with known
hypersensitivity to fenbendazole, pyrantel,
praziquantel or any of the excipients should
administer the veterinary medicinal product
with caution.

When handling the veterinary medicinal
product (especially in breeding facilities with a
larger number of dogs), personal protective
equipment consisting of rubber or latex gloves
should be used.

Do not eat, drink or smoke while handling the
product.

Wash your hands with water and soap after
using the product.

Echinococcosis poses a risk to humans and is
notifiable to the World Organization for Animal
Health (WOAH). In case of occurrence of
echinococcosis, special instructions for
treatment, monitoring and ensuring personal
safety must be followed. Consult
parasitologists or parasitological institutes.
Poultry aroma with yeast contains poultry
protein.

Pregnancy:

Dosing should be strictly observed in pregnant
bitches.

Interaction with other medicinal products and other forms of interaction:

Do not administer concurrently with piperazine-
based products, or other antiparasitics.

Overdose (symptoms, emergency procedures, antidotes)

A tenfold dose has no side effects on the target
animals.

13. SPECIAL PRECAUTIONS FOR THE DISPOSAL OF UNUSED PRODUCTS OR WASTE MATERIALS, IF ANY

Any unused veterinary medicinal product or
waste materials derived from such veterinary
medicinal product should be disposed of in
accordance with local requirements.
Medicines should not be disposed of via
wastewater or household waste.

14. DATE ON WHICH THE PACKAGE LEAFLET WAS LAST APPROVED

22/07/2024

15. OTHER INFORMATION

For animal treatment only.

Veterinary medicinal product not subject to
prescription.

For any information about this veterinary
medicinal product, please contact the local
representative of the marketing authorisation
holder.



Dimensions (mm):

Colours:

Prepared by:

Date:

A6 (105 × 148 mm) duplex printing

black

Hanáková

20. 01. 2025

Approved by:

Date:

Monika

Hladká

Digitálně

podepsal

Monika Hladká

Datum:

2025.01.29

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minimum space for reprinting batch and expiration

11x5 mm

Dimensions (mm): 60x30
Colours: Black

Prepared by: Raška
Date: 27. 10. 2021



CANIVERM

oral paste

Fenbendazole, pyrantel embonate, praziquantel

1 ml of the paste contains:

Fenbendazole	75 mg
Pyrantel embonate	72 mg
Praziquantel	25 mg

4 ml

Route(s) of administration: Oral.

Withdrawal period:

Not intended for food-producing animals.

**Shelf life after first opening
the container:** 6 months.

Once opened use by:

For animal treatment only.



Bioveta, a. s.
Czech Republic



Batch:
EXP:

319/E/4ml/GB0/3

minimální prostor pro dotisk šarže a expirace
minimum space for reprinting batch and expiration

11×5 mm

Dimensions (mm): 50×50
Colours: CMYK

Prepared by: Raška
Date: 12. 4. 2023

