

Leaflet – Fleas serve as intermediate hosts for a common type of tapeworm – *Dipylidium caninum*. Tapeworm infestation is certain to recur unless control of intermediate hosts such as fleas, mice, etc. is undertaken.

Dispensing this medicine requires a veterinary surgeon's prescription
For use in animals only

1. Name, form, and strength of the veterinary medicinal product

Prazitel Cat veterinary tablets

2. Active ingredient

Each tablet contains:

Praziquantel 20 mg, Pyrantel embonate 230 mg.

Inactive ingredients and allergens: See Section 13 'Additional Information'.

3. What is this medicine intended for?

For treating infections caused by tape worms and round worms in cats.

Therapeutic group:

Anthelmintic.

4. Contraindications

Do not use concomitantly with products containing piperazine.

Do not use in kittens under 6 weeks old.

Do not use in animals with known hypersensitivity to the active substances or to any of the excipients.

Do not use in pregnancy.

5. Side effects

None known.

If you notice any serious side effects, please inform your veterinary surgeon.

You can report side effects to the Ministry of Health (MoH) by following the link 'Reporting Side Effects of Medication' on the MoH home page (www.health.gov.il) which links to an online form for reporting side effects. You can also use this link:

<https://forms.gov.il/globaldata/getsequence/getsequence.aspx?formType=AdversEffectMedic@moh.gov.il>

6. Target species

Cats

7. Route of administration and dosage in the target animal

Single dose: for oral administration.

The recommended dose is: 20 mg pyrantel (equivalent to 57.5 mg/kg pyrantel embonate) and 5 mg/kg praziquantel. This is equivalent to 1 tablet per 4 kg body weight.

Dosage table:

Body weight (kg)	Tablets
1.0-2.0	1/2
2.1-4.0	1
4.1-6.0	1½
6.1-8.0	2

The tablet should be given to the cat directly, but if necessary can be disguised in food.

8. How to use the medicinal product

To ensure administration of a correct dose, body weight should be determined as accurately as possible.

9. Withdrawal period

N/A.

10. Precautions

• Special precautions for use in the target animal

Parasitic resistance to a certain class of anthelmintics can occur after frequent and repeated use of an anthelmintic from this class.

Fleas serve as intermediate hosts for a common type of tapeworm – *Dipylidium caninum*. Tapeworm infestation is certain to recur unless control of intermediate hosts such as fleas, mice, etc. is undertaken.

• Special precautions to be taken by the person administering the medicinal product

In case of accidental ingestion, seek medical advice and show the package leaflet to the physician.

In the interests of good hygiene, persons administering the tablets directly to a cat or adding them to the cat's food should wash their hands afterwards.

For animal treatment only.

• Pregnancy and lactation

Do not use during pregnancy; may be used during lactation.

• Drug interactions and other interactions

Do not use this medicinal product concomitantly with products containing piperazine.

• Overdose

After doses higher than 5 times the recommended dose, signs of intolerance such as vomiting have been observed.

11. Storage instructions

• Prevent poisoning! To prevent poisoning, keep this, and all other medicines, in a closed place out of sight and reach of children and/or infants.

• Do not use after the expiry date (Exp. Date) stated on the package. The expiry date refers to the last day of that month.

• Do not store above 25°C.

• Do not remove the tablets from the package until ready to use them.

12. Instructions for destroying the medicinal product/ any remaining medicinal product after use

Discard any unused split tablets immediately.

Any unused veterinarian medicinal preparation or waste from such veterinarian medicinal preparations must be disposed of according to the local regulatory requirements.

Do not discard medicines via wastewater or household waste.

Ask your veterinary surgeon how to dispose of medicines no longer required. These measures should help to protect the environment.

13. Other information

• In addition to the active ingredients the medicine also contains: Maize starch, Microcrystalline Cellulose, Croscopovidone, Magnesium Stearate, Colloidal Anhydrous Silica, Opadry complete film coating System 03F28415 White, Grilled meat flavor.

• What the medicine looks like and what are the contents of the package:

White to off-white round, biconvex coated tablets with a score line on one side and plain on the other side.

Available in packages of 104 tablets.

Registration holder: E.L. Medi Market Ltd. 18 Hakadar St., Netanya 42138.

Manufacturer name: Chanelle Pharmaceuticals Manufacturing Ltd., Loughrea, Co.Galway, Ireland.

This leaflet was reviewed and approved by the Ministry of Health in: February 2017.

Registration number of the medicine in the National Drug Registry of the Ministry of Health: 157-98-34522-00.