

CONSUMER PACKAGE INSERT FOR A VETERINARY PREPARATION

The medicine is dispensed with a veterinarian's prescription only
For animal use only

1. NAME OF THE VETERINARY MEDICINE, ITS FORM AND STRENGTH

Prazitel Cat

Veterinary Tablets

2. ACTIVE INGREDIENTS

Each tablet contains:

Praziquantel 20 mg, Pyrantel Embonate 230 mg.

Inactive ingredients and allergens in the preparation: see section 13 "Further Information".

3. WHAT IS THE MEDICINE INTENDED FOR

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

Therapeutic group:

Anthelmintic.

4. CONTRAINDICATIONS

Do not use the preparation together with preparations containing piperazine.

Do not use in kittens less than six weeks of age.

Do not use the preparation for treatment in animals with known sensitivity to the active ingredients or to any one of the inactive ingredients.

Do not use this preparation during pregnancy.

5. SIDE EFFECTS

In very rare cases, slight and temporary disturbances in the digestive system such as excessive salivation and/or vomiting, as well as temporary neurological disturbances such as ataxia (lack of limb co-ordination), may appear. If you notice any side effects, even those not appearing in the leaflet, or if you think that the medicine is somewhat ineffective, please inform the veterinary doctor.

Side effects can be reported to the Ministry of Health by clicking on the link "Report Side Effects of Drug Treatment" found on the Ministry of Health homepage (www.health.gov.il) that directs you to the online form for reporting side effects, or by entering the link:

<https://sideeffects.health.gov.il>

6. TARGET SPECIES

Cats.

7. DOSAGE AND MODE OF ADMINISTRATION

A single dose: for oral administration.

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

Dosage table:

Body weight (kg)	Tablets
1.0-2.0	½
2.1-4.0	1
4.1-6.0	1½
6.1-8.0	2

The tablets should be given to the cat directly, but if necessary, they can be dispersed in food.

8. HOW TO USE THE PREPARATION

To ensure administration of an appropriate dose, body weight should be determined as accurately as possible.

9. WITHDRAWAL PERIOD

Not applicable.

10. WARNINGS

- Special warnings regarding use of the medicine in treatment of the target animal

The parasites can develop resistance to a particular class of anthelmintics following repeated and frequent use of an anthelmintic from this class.

Fleas serve as intermediate hosts for a common type of tapeworms –

Dipylidium caninum. The chance for tapeworm infestation is certain to recur unless control of intermediate hosts, such as fleas, mice, and the like is undertaken.

- Special warnings regarding safety in the use of the medicine in animals
Since the tablets have a flavor, they must be stored in a safe place out of the reach of animals.

- Special warnings regarding the safety of the person handling the preparation
In case of accidental ingestion, seek medical assistance and show the package leaflet to the doctor.

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 in the treated animal

Do not use this preparation during pregnancy but it may be used during the lactation period.

- Reactions with other drugs and other types of interactions

Do not use the preparation together with preparations containing piperazine compounds.

- Overdosage

After taking a dose five times greater than the recommended dosage, signs of intolerance such as vomiting, for example, have been observed.

11. STORAGE INSTRUCTIONS

- Avoid poisoning! This medicine and any other medicine should be kept in a safe place out of the reach and sight of children and/or infants in order to avoid poisoning.

- Do not use the medicine after the expiry date (exp. date) that appears 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 their use.

12. INSTRUCTIONS REGARDING DISPOSAL OF THE PREPARATION/ REMNANTS OF THE PREPARATION AFTER USE

Discard any unused half tablets immediately.

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

Do not dispose of medicines via the household wastewater or household waste.

Ask your veterinary doctor how to dispose of medicines no longer required. This will help to protect the environment.

13. FURTHER INFORMATION

- In addition to the active ingredients the medicine also contains:

Maize starch, Microcrystalline cellulose, Crospovidone, Magnesium stearate, Colloidal anhydrous silica, Opadry complete film coating system 03F28415 white, Grilled meat flavour.

- What does the medicine look like and what is the content of the package:
Film-coated, white-cream-colored, round tablets, with a score line on one side and plain on the other side.

Marketed in a package of 104 tablets.

License Holder and its Address:

A.L.Medi-Market Ltd.,
3 Hakatif St., Emek Hefer Industrial Park,
3877701.

Name of Manufacturer:

Chanelle Pharmaceuticals Manufacturing Ltd., Loughrea, Co. Galway, Ireland

Registration number of the medicine in the National Drug Registry of the Ministry of Health:

157-98-34522-00

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