CONSUMER PACKAGE INSERT FOR VETERINARY MEDICINAL PRODUCT

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

1. NAME OF THE MEDICINAL PRODUCT:
Panacur 4 % Veterinary

2. COMPOSITION:
Active ingredient:
1 g powder contains:
Fenbendazole - 40 mg

The list of excipients is detailed in section 13.

3. WHAT IS THE MEDICINAL PRODUCT INTENDED FOR:
For the treatment of infestation of pigs with gastrointestinal worms in their immature and mature stages, kidney worms and/or lungworms and, for example:
red stomach worms, nodular worms, roundworms, whipworms, kidney worms and lungworms.

4. CONTRAINDICATIONS:
Do not use in case of known sensitivity to the active ingredient or one of the other ingredients.

5. SIDE EFFECTS:

None known.

If you notice side effects or any other reactions not mentioned in this package insert, please notify your veterinarian.

Side effects can be reported to the Ministry of Health by clicking on the link "Reporting Side Effects due to 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 ANIMALS:
Pigs.

7. METHOD OF ADMINISTRATION AND DOSAGE:
Powder for the preparation of medicated feed

The preparation can be offered with the usual feed and consumed separately or used to prepare a medicated feed (mixed with industrial feed or an appropriate commercially available compound feed).

Unless otherwise prescribed, adhere exactly to the dose.

Recommended dose: 5 mg Fenbendazole per kg body weight.

For infestation with swine kidney worms (Stephanurus dentatus): 10 mg Fenbendazole per kg body weight.

1. Single treatment with the therapeutic dose

Single treatment
A graduated measuring beaker is included for 500 g and 2.5 kg doses.

Mass treatment
In order to prepare a medicated feed, the therapeutic dose (5 mg Fenbendazole per kg body weight) should be mixed into the daily ration.
For example, the following procedure should be used:

Animal species	Feed consumption	Powder per tonne of compound feed
Fattening pigs - Approx. 20 kg body weight	1 kg/day/animal	2.5 kg
Sows - Approx. 200 kg body weight	2 kg/day/animal (basic ration)	12.5 kg

For infections with kidney worms, the dose should be doubled (10 mg Fenbendazole per kg body weight).
For attacks with whip worms and/or lungworms, treatment should be administered according the procedure described in Section 2.

2. Distribute the therapeutic dose across 5-15 days
When preparing a medicated feed, the therapeutic dose (5 mg Fenbendazole per kg body weight) should be mixed into the compound feed ration intended for 5-15 days.
For example, the following procedure should be used:

Animal species	Treatment duration	Feed consumption	Powder per tonne of compound feed
Fattening pigs - approx. 20 kg body weight	5 days 10 days 15 days	1 kg/Day/Animal	0.5 kg 0.25 kg 0.17 kg
Sows - approx. 200 kg body weight	5 days 10 days 15 days	2 kg/day/animal (Basic ration)	2.5 kg 1.25 kg 0.83 kg

In order to prepare pre-mixes (2 kg per tonne feed), wheat middlings are recommended.
The powder is odourless, tasteless and accepted by the animals.

8. HOW TO USE THE PRODUCT:

The preparation can also be administered to animals that are seriously ill, pregnant or generally in poor condition.
Dietary measures are not required before or after treatment. Repeat the treatment after reinfection.

9. WITHDRAWAL PERIOD:

Meat and offal: 7 days

10. WARNINGS:

- Special warnings
  None.

- Special warnings relating to use of the medicine in animals
  Not applicable.

- Special warnings relating to the safety of the person administering the medicinal product
  Not applicable.

- Pregnancy and lactation in treated animal
  Can be used during pregnancy.

- Interactions with other medicinal products and other types of interactions
  None known.

- Overdose
  No information

- Incompatibility
  None known.

11. STORAGE INSTRUCTIONS:

- Avoid poisoning! This medicinal product and any other medicinal products, should be kept in a closed place, out of the reach and sight of children and/or infants in order to protect from poisoning.
- "Do not use the medicinal product after the expiry date (exp. Date) appearing on the package. The expiry date refers to the last day of that month".
- Store below 25^{o}C.
- Store in dry place.
- Protect from light.
- Do not store in refrigerator or freezer.

- Keep in the original packaging.

12. INSTRUCTIONS REGARDING DISPOSAL OF THE MEDICINAL PRODUCT/REMNANTS OF THE MEDICINAL PRODUCT AFTER USE:

Any unused veterinary medicinal product or waste materials derived from such veterinary medicinal product should be disposed of as toxic waste, do not throw to sewer.

13. ADDITIONAL INFORMATION:

- In addition to the active substance(s), the medicine also contains:
  Lactose monohydrate
  Calcium carbonate
  Maize starch
  Colloidal anhydrous silica

- Package size
  2.5 kg

- What the medicine looks like and the content of the package:
  White to grayish powder.
  Plastic bucket (polypropylene) in white color with polyethylene inner bag and polypropylene measuring beaker.

- Registration holder and its address:
  Intervet Israel Ltd., Industrial Park Neve Ne'eman, Hod Hasharon 45240.

- Manufacturer name and address:
  Intervet GesmbH, Siemensstrasse 107, 1210 Vienna, Austria

- This leaflet was checked and approved by the Ministry of Health in January 2020.

- **Registration number of the medicinal product in the National Drug Registry of the Ministry of Health: 163-44-35216-00**