

PARTICULARS TO APPEAR ON THE OUTER PACKAGE
Cardboard box 10 x 10 x 10.2 g

1. NAME OF THE VETERINARY MEDICINAL PRODUCT

Panacur Equine 222 mg/g Granules

2. STATEMENT OF ACTIVE SUBSTANCES

Each gram contains 222 mg fenbendazole.

3. PACKAGE SIZE

10 boxes containing 10 sachets of 10.2 g

4. TARGET SPECIES

Horses and other equines.

5. INDICATIONS

Horse wormer.

6. ROUTES OF ADMINISTRATION

Oral use.

Routine treatment: One sachet per 300 kg bodyweight as a single dose.

Control of migrating large redworm larvae and encysted mucosal small redworm:
One sachet per 300 kg bodyweight daily for five consecutive days.

Sprinkle granules onto the horse's grain or concentrate feed with the full daily dosage given in one feed.

Treatment should be repeated when natural reinfestation with worms occurs.

7. WITHDRAWAL PERIODS

Withdrawal periods:

Not to be used in horses intended for human consumption.

Treated horses may never be slaughtered for human consumption.

The horse must have been declared as not intended for human consumption under national horse passport legislation.

8. EXPIRY DATE

Exp. {mm/yyyy}

9. SPECIAL STORAGE PRECAUTIONS

This veterinary medicinal product does not require any special storage conditions. Keep the sachets in the outer carton.

10. THE WORDS “READ THE PACKAGE LEAFLET BEFORE USE”

Read the package leaflet before use.

11. THE WORDS “FOR ANIMAL TREATMENT ONLY”

For animal treatment only.

12. THE WORDS “KEEP OUT OF THE SIGHT AND REACH OF CHILDREN”

Keep out of the sight and reach of children.

13. NAME OF THE MARKETING AUTHORISATION HOLDER

Marketing Authorisation Holder:
Intervet International B.V.

14. MARKETING AUTHORISATION NUMBERS

Vm 06376/4092

15. BATCH NUMBER

Lot {number}

Class of anthelmintic:

1-BZ

PARTICULARS TO APPEAR ON THE OUTER PACKAGE
Cardboard box 10 x 10.2 g sachets

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8. EXPIRY DATE

Exp. {mm/yyyy}

9. SPECIAL STORAGE PRECAUTIONS

This veterinary medicinal product does not require any special storage conditions.
Keep the sachets in the outer carton.
When sold individually, sachets should be accompanied by a leaflet.

10. THE WORDS “READ THE PACKAGE LEAFLET BEFORE USE”

Read the package leaflet before use.

11. THE WORDS “FOR ANIMAL TREATMENT ONLY”

For animal treatment only.

12. THE WORDS “KEEP OUT OF THE SIGHT AND REACH OF CHILDREN”

Keep out of the sight and reach of children.

13. NAME OF THE MARKETING AUTHORISATION HOLDER

Marketing Authorisation Holder:
Intervet International B.V.

14. MARKETING AUTHORISATION NUMBERS

Vm 06376/4092

15. BATCH NUMBER

Lot {number}

Class of anthelmintic:

1-BZ

PARTICULARS TO APPEAR ON THE IMMEDIATE PACKAGE
10.2 g sachet

1. NAME OF THE VETERINARY MEDICINAL PRODUCT

Panacur Equine 222 mg/g Granules

2. STATEMENT OF ACTIVE SUBSTANCES

Each gram contains 222 g fenbendazole.

10.2 g sachet

3. TARGET SPECIES

Horses and other equines.

4. ROUTES OF ADMINISTRATION

Oral use.

Read the package leaflet before use.

5. WITHDRAWAL PERIODS

Withdrawal periods:

Not to be used in horses intended for human consumption.

Treated horses may never be slaughtered for human consumption.

The horse must have been declared as not intended for human consumption under national horse passport legislation.

6. EXPIRY DATE

Exp. {mm/yyyy}

7. SPECIAL STORAGE PRECAUTIONS

This veterinary medicinal product does not require any special storage conditions.
Keep the sachets in the outer carton.

8. NAME OF THE MARKETING AUTHORISATION HOLDER

Intervet International B.V.

9. BATCH NUMBER

Lot {number}

PARTICULARS TO APPEAR ON THE PACKAGE LEAFLET

PACKAGE LEAFLET

1. Name of the veterinary medicinal product

Panacur Equine 222 mg/g Granules

2. Composition

Each gram contains 222mg fenbendazole.

A white to yellowish-white granular powder.

3. Target species

Horses and other equines.

4. Indications for use

A broad spectrum anthelmintic for the treatment and control of adult and immature roundworms of the gastrointestinal tract in horses and other equines.

The veterinary medicinal product effectively treats and controls the following roundworm infections:

Large strongyles (adults and migrating larval stages of *S. vulgaris*; adults and tissue larval stages of *S. edentatus*).

Benzimidazole susceptible adult and immature small strongyles (cyathostomes) including encysted mucosal 3rd and 4th stage larvae and it is also effective against inhibited 3rd stage larvae (encysted) in the mucosa.

Adult and immature *Oxyuris* spp., *Strongyloides* spp. and *Parascaris equorum*.

This veterinary medicinal product has an ovicidal effect on nematode eggs.

5. Contraindications

None known.

6. Special warnings

Special warnings:

Care should be taken to avoid the following practices because they increase the risk of development of resistance and could ultimately result in ineffective therapy:

- Too frequent and repeated use of anthelmintics from the same class, over an extended period of time.
- Under dosing, which may be due to underestimation of body weight, misadministration of the veterinary medicinal product, or lack of calibration of the

dosing device (if any).

Suspected clinical cases of resistance to anthelmintics should be further investigated using appropriate tests (e.g. Faecal Egg Count Reduction Test). Where the results of the test(s) strongly suggest resistance to a particular anthelmintic, an anthelmintic belonging to another pharmacological class and having a different mode of action should be used.

Resistance to fenbendazole has been reported in cyathostomes in horses. Therefore, the use of this veterinary medicinal product should be based on local (regional, farm) epidemiological information about susceptibility of nematodes and recommendations on how to limit further selection for resistance to anthelmintics.

Special precautions for safe use in the target species:

Not applicable.

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

Direct contact with the skin should be avoided. Avoid inhalation of granule dust. Wash hands after use.

Pregnancy:

Pregnant mares and young foals may be safely treated with fenbendazole at therapeutic dosage levels.

Interaction with other medicinal products and other forms of interaction:

None known.

Overdose:

Benzimidazoles are unlikely to cause any reactions in the target species.

Major incompatibilities:

None known.

7. Adverse events

Horses and other equines.

None known.

Reporting adverse events is important. It allows continuous safety monitoring of a product. 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 contact, in the first instance, your veterinarian. You can also report any adverse events to the marketing authorisation holder or the local representative of the marketing authorisation holder using the contact details at the end of this leaflet, or via your national reporting system at:

Website: <https://www.gov.uk/report-veterinary-medicine-problem/animal-reacts-medicine>

e-mail: adverse.events@vmd.gov.uk

8. Dosage for each species, routes and method of administration

Oral use.

The veterinary medicinal product is odourless and tasteless and should be mixed with your horse's concentrate or grain feed with the full daily dosage given in one feed. It is not necessary to withhold feed before or after treatment.

Dosage and administration

Routine treatment: Administer orally 5 g of the veterinary medicinal product per 150 kg bodyweight.
(= 7.5 mg fenbendazole/kg bodyweight).

Each sachet contains 10.2 g granules and can be used as follows:

Foals and ponies up to 300 kg bodyweight	1 sachet
Thoroughbreds and other breeds of horses up to 600 kg bodyweight	2 sachets
Heavy hunters, heavy draft horses	3 sachets
Donkeys	1 sachet

Increased dosing for specific infections

Five-day course:

For the treatment and control of migrating larval stages of large strongyles and encysted mucosal 3rd and 4th stage larvae and inhibited 3rd stage small strongyle larvae (encysted) in the mucosa, administer 5 g of the veterinary medicinal product per 150 kg bodyweight daily for 5 consecutive days.

(= 7.5 mg fenbendazole/kg bodyweight daily for 5 days)

Single dose treatments:

For the treatment and control of encysted mucosal stages of small strongyles, administer 20 g of the veterinary medicinal product per 150 kg bodyweight.

(= 30 mg fenbendazole/kg bodyweight)

For the treatment and control of migrating stages of large strongyles, administer 40 g of the veterinary medicinal product per 150 kg bodyweight.

(= 60 mg fenbendazole/kg bodyweight)

Diarrhoea caused by *Strongyloides westeri* in two- to three-week-old suckling foals should be treated with Panacur 10% Suspension at a dose rate of 25 ml per 50 kg bodyweight.

(= 50 mg fenbendazole/kg bodyweight)

Recommended dosing programme

Inappropriate use of anthelmintics may increase resistance selection pressure and lead to reduced efficacy. The decision to use the product should be based on professional advice and take into account current best practice recommendations for parasite control.

Seek veterinary advice for appropriate monitoring, stock management and dosing programmes to allow optimum endoparasite control.

9. Advice on correct administration

To ensure a correct dosage, bodyweight should be determined as accurately as possible. The use of a weigh band is recommended.

The use of suitably calibrated measuring equipment is recommended. Accuracy of the dosing device should be checked.

10. Withdrawal periods

Not to be used in horses intended for human consumption.

Treated horses may never be slaughtered for human consumption.

The horse must have been declared as not intended for human consumption under national horse passport legislation.

11. Special storage precautions

Keep out of the sight and reach of children.

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

Keep the sachets in the outer carton.

12. Special precautions for disposal

Medicines should not be disposed of via wastewater.

The veterinary medicinal product should not enter water courses as fenbendazole may be dangerous for fish and other aquatic organisms. Do not contaminate ponds, waterways or ditches with the veterinary medicinal product or used container.

Use take-back schemes for the disposal of any unused veterinary medicinal product or waste materials derived thereof in accordance with local requirements and with any applicable national collection systems. These measures should help to protect the environment.

Ask your veterinary surgeon or pharmacist how to dispose of medicines no longer required.

13. Classification of veterinary medicinal products

Veterinary medicinal product subject to prescription.

14. Marketing Authorisation Numbers and Pack Sizes

Vm 06376/4092

Pack sizes:

10 x 10.2 g sachets.

100 x 10.2 g sachets.

Not all pack sizes may be marketed.

15. PID LINK (Do not print heading)

[The following statement must be included where reference to the European Union Product Database is included on the product information. This statement is relevant to both UK(GB) and UK(NI) products:]

Find more product information by searching for the 'Product Information Database' on www.gov.uk.

16. Contact details

Marketing authorisation holder:

Intervet International B.V.
Wim de Körverstraat 35
5831 AN Boxmeer
Netherlands

Manufacturer responsible for batch release:

Intervet GesmbH
Siemenstrasse 107
1210 Vienna
Austria

Local representative:

MSD Animal Health UK Limited
Walton Manor, Walton
Milton Keynes
MK7 7AJ, United Kingdom

Contact details to report suspected adverse reactions:

UK(GB)

MSD Animal Health UK Ltd.
Tel.: +44 (0)1908 685685

UK(NI)

Intervet Ireland Ltd.
Tel.: +353 (0)1 2970220

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

17. Other information

POM-VPS

Environmental properties

Fenbendazole is toxic to fish and other aquatic organisms.

Class of anthelmintic:

1-BZ

Gavin Hall

Approved 22 February 2026