

PARTICULARS TO APPEAR ON THE OUTER PACKAGE
DISPLAY CARTON

1. NAME OF THE VETERINARY MEDICINAL PRODUCT

Strongid-P Oral Paste 439 mg/g

2. STATEMENT OF ACTIVE SUBSTANCES

Pyrantel Embonate 439 mg/g

3. PACKAGE SIZE

10 oral syringes

4. TARGET SPECIES

For horses, ponies, donkeys and foals over four weeks of age.

5. INDICATIONS

6. ROUTES OF ADMINISTRATION

See oral syringe carton for full directions for use.

7. WITHDRAWAL PERIODS

Withdrawal period: Meat and offal - zero days.

8. EXPIRY DATE

Exp. {mm/yyyy}

9. SPECIAL STORAGE PRECAUTIONS

Do not store above 25°C. Protect from direct sunlight.
Store in tightly closed original container

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

Read the label 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

Elanco GmbH

14. MARKETING AUTHORISATION NUMBERS

Vm 52127/5081

Vm 52127/3016

15. BATCH NUMBER

Lot {number}

PARTICULARS TO APPEAR ON THE IMMEDIATE PACKAGE
SYRINGE

1. NAME OF THE VETERINARY MEDICINAL PRODUCT

Strongid-P Oral Paste 439 mg/g

2. STATEMENT OF ACTIVE SUBSTANCES

Pyrantel Embonate 439 mg/g

3. TARGET SPECIES

For horses, ponies, donkeys and foals over four weeks of age.

4. ROUTES OF ADMINISTRATION

See oral syringe carton for full directions for use.

5. WITHDRAWAL PERIODS

Withdrawal period: Meat and offal - zero days.

6. EXPIRY DATE

Exp. {mm/yyyy}

After first opening, use by:
Once opened use within 2 months.

7. SPECIAL STORAGE PRECAUTIONS

Do not store above 25°C and protect from direct sunlight.
Store in tightly closed original container.

8. NAME OF THE MARKETING AUTHORISATION HOLDER

Elanco GmbH

9. BATCH NUMBER

Lot {number}

PARTICULARS TO APPEAR ON THE IMMEDIATE PACKAGING WHERE THERE IS NO PACKAGE LEAFLET, i.e. Combined label and package leaflet CARTON

1. NAME OF THE VETERINARY MEDICINAL PRODUCT

Strongid-P Oral Paste 439 mg/g

2. COMPOSITION

Pyrantel Embonate 439 mg/g

A smooth, pale yellow to buff paste.

3. PACKAGE SIZE

26 g

4. TARGET SPECIES

For horses, ponies, donkeys and foals over four weeks of age.

5. INDICATIONS FOR USE

Indications for use

The veterinary medicinal product is effective against all the important gut-dwelling roundworms found in the horse, and also tapeworms when twice the normal dose level is used.

The veterinary medicinal product is effective against benzimidazole-resistant strains of small redworms.

The broad-spectrum activity includes:

Large Roundworms

Parascaris equorum.

Large Redworms

(large strongyles) *Strongylus vulgaris*, *Strongylus edentatus*, *Strongylus equinus* and *Triodontophorus* spp.

Small Redworms

(small strongyles including benzimidazole-resistant strains) *Cyathostomum* spp.

Seatworms/pinworms *Oxyuris equi*.

Tapeworms *Anoplocephala perfoliata*.

6. CONTRAINDICATIONS

Contraindications

Do not use in cases of hypersensitivity to the active substance or to any of the excipients.

7. 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.
- Underdosing, which may be due to underestimation of body weight, misadministration of the 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 anthelmintics belonging to another pharmacological class and having a different mode of action should be used.

Resistance to pyrantel has been reported in cyathostomes in horses (also widespread in the US and Canada). Therefore the use of this product should be based on a 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:

The same oral syringe should only be used to dose two animals if they are both healthy and are either running together or are on the same premises and in direct contact with each other.

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

Avoid contact with skin. Wash hands and any other parts of the body which come into contact with the product.

Special precautions for the protection of the environment:

Not applicable.

Pregnancy and lactation:

The veterinary medicinal product is specifically recommended for use in mares which may be pregnant and/or lactating.

Interactions with other medicinal products and other forms of interaction:

Combined administration of pyrantel and levamisole or piperazine is not recommended.

Overdose:

The veterinary medicinal product has an extremely wide safety margin and overdosage should not produce any adverse reactions.

Major incompatibilities:

None known.

8. ADVERSE EVENTS

Adverse events

Horses, ponies, donkeys, foals:

Very rare (<1 animal / 10,000 animals treated, including isolated reports):

Colic ¹ , Intestinal impaction ²
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¹May be seen within 30 minutes of treatment.

²May occur in foals infected with high numbers of *Parascaris equorum*.

Reporting adverse events is important. It allows continuous safety monitoring of a product. If you notice any side effects, even those not already listed on this label, 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 using the contact details on this label, 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

9. DOSAGE FOR EACH TARGET SPECIES, ROUTES AND METHOD OF ADMINISTRATION

Dosage for each species, routes and method of administration

Administration

The veterinary medicinal product is recommended for direct oral administration in horses, ponies, donkeys and foals over four weeks of age. It is not necessary to withhold any feed prior to administration or change normal exercise routines following dosing.

The following method of administration is recommended:

1. Position the locking ring over the appropriate mark on the plunger.
2. Remove the cap from the nozzle.

3. The paste is best deposited on the upper surface of the tongue. Introduce the nozzle end of the oral syringe at the corner of the mouth. Direct the oral syringe backwards and depress the plunger to deposit the paste onto the tongue. Providing the paste is given in this way it is unlikely that any rejection will occur. Raising of the horse's head by a hand under the chin sometimes helps with swallowing.

4. Store in tightly closed original container. Any product remaining 2 months after opening should be discarded.

5. The same oral syringe should only be used to dose two animals if they are both healthy and are either running together or are in direct contact with each other.

Dosage: For the control and treatment of adult infections of large and small redworm, including benzimidazole-resistant strains, the large roundworm and pinworm or seatworm.

Dosage to include the control of tapeworms:

To include the control of tapeworm the dose should be doubled.

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.

DOSE TABLE FOR CONTROL OF ROUNDWORMS

N.B. TO INCLUDE CONTROL OF TAPEWORM PLEASE DOUBLE THE USUAL ROUNDWORM DOSE

To ensure administration of a correct dose, body weight should be determined as accurately as possible; accuracy of the dosing device should be checked.

Class of Horse	Height in Hands	Weight in Kg	Dose In Whole Oral Syringes
Heavy Hunters	16+	600-700	1.25 oral syringes
Thoroughbreds, Hunters Hacks, Arabs	14.2-16.2	450-600	1 oral syringe
Ponies	12.2-14.2	300-450	0.75 oral syringe
Small Ponies and Donkeys	9.0-12.2	150-300	0.5 oral syringe
Miniature Ponies	Less than 9.0	Up to 150 kg	0.25 oral syringe

10. ADVICE ON CORRECT ADMINISTRATION

Advice on correct administration

11. WITHDRAWAL PERIODS

Withdrawal periods

Meat and offal - zero days.

12. SPECIAL STORAGE PRECAUTIONS

Special storage precautions

Keep out of the sight and reach of children.

Do not store above 25°C. Protect from direct sunlight.

Store in tightly closed original container.

Do not use this veterinary medicinal product after the expiry date which is stated on the carton after Exp. The expiry date refers to the last day of that month.

13. SPECIAL PRECAUTIONS FOR DISPOSAL

Special precautions for disposal

Medicines should not be disposed of via wastewater.

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.

14. CLASSIFICATION OF VETERINARY MEDICINAL PRODUCTS

Classification of veterinary medicinal products

Veterinary medicinal product subject to prescription.

15. MARKETING AUTHORISATION NUMBERS AND PACK SIZES

Vm 52127/5081

Vm 52127/3016

Pack sizes

Carton with 10 syringes containing 26 g of product

16. 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.

17. CONTACT DETAILS

Contact details

Marketing authorisation holder and contact details to report suspected adverse reactions:

Elanco GmbH
Heinz-Lohmann Strasse 4
Groden
27472 Cuxhaven
Germany

Tel: +44 3308221732

PV.GBR@elancoah.com

Manufacturer responsible for batch release:

Purna Pharmaceuticals NV,
Rijksweg 17,
Puurs,
B-2870,
Belgium

18. OTHER INFORMATION

POM-VPS

19. THE WORDS “FOR ANIMAL TREATMENT ONLY”

For animal treatment only.

20. EXPIRY DATE

Exp {mm/yyyy}

Once opened use within 2 months.

21. BATCH NUMBER

Lot {number}

Gavin Hall

Approved: 24 February 2026