PARTICULARS TO APPEAR ON THE OUTER PACKAGE – Carton box Flat-bottle

1. NAME OF THE VETERINARY MEDICINAL PRODUCT

Panacur 10% oral suspension

2. STATEMENT OF ACTIVE SUBSTANCES

Each ml contains:
Fenbendazole 100 mg

3. PACKAGE SIZE

1 Litre
2 Litres
5 Litres
10 Litres

1 Litre
1000 doses for 20 kg lambs, 400 doses for 50 kg ewes
66 doses for 200 kg cattle, 22 doses for 600 kg horses

2 Litre
2000 doses for 20 kg lambs, 800 doses for 50 kg ewes
133 doses for 200 kg cattle, 44 doses for 600 kg horses

5 Litre
5000 doses for 20 kg lambs, 2000 doses for 50 kg ewes
333 doses for 200 kg cattle, 111 doses for 600 kg horses

10 Litre
10000 doses for 20 kg lambs, 4000 doses for 50 kg ewes
666 doses for 200 kg cattle, 222 doses for 600 kg horses

4. TARGET SPECIES

Cattle, sheep, horses and other equines.

5. INDICATIONS

For the treatment and control of gastro-intestinal roundworm infections in cattle, sheep and horses and of lungworm and tapeworm infections in cattle and sheep.

6. ROUTES OF ADMINISTRATION

Oral use.
Shake container before use.
Cattle and Horses: 1 ml per 13 kg bodyweight (7.5 mg fenbendazole per kg bodyweight) to be given orally (see table). For horses, mix the product with grain or concentrate feed and give the full dose as one administration. Pregnant mares and foals may be treated safely with fenbendazole at therapeutic levels. For the treatment of migrating larval and tissue stages of large strongyles and encysted stages of small strongyles in horses, the dose should be repeated daily for five days.
Sheep: 0.5 ml per 10 kg bodyweight (5 mg fenbendazole per kg bodyweight) to be given orally (see table).
A dosing scheme per target species as well as guidance regarding the number of doses per pack size is provided on the carton box.
To ensure a correct dosage, bodyweight should be determined as accurately as possible. Accuracy of the dosing device should be thoroughly checked.
Treatment should be repeated when reinfestation occurs.

7. WITHDRAWAL PERIODS

Withdrawal periods:
Cattle:
Meat and offal: 12 days
Milk: 5 days

Sheep:
Meat and offal: 15 days
Milk: 7 days

Horses:
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}
Liquid feed containing the product will remain stable for up to 3 months.

9. SPECIAL STORAGE PRECAUTIONS

Do not store above 25 °C.
Do not freeze.
Protect from frost.

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

MA Holder:
Intervet International B.V.

Distributor in Northern Ireland:
Intervet Ireland Ltd.

14. MARKETING AUTHORISATION NUMBERS

Vm 06376/4076

15. BATCH NUMBER

Lot {number}

Class of anthelmintic: (1-BZ)

PARTICULARS TO APPEAR ON THE IMMEDIATE PACKAGING WHERE THERE IS NO PACKAGE LEAFLET, Front and back label - Flat-bottle

1. NAME OF THE VETERINARY MEDICINAL PRODUCT

Panacur 10% oral suspension

2. COMPOSITION

Each ml contains:

Active substance:

Fenbendazole	100 mg

Excipients:

Benzyl alcohol	4.835 mg
Sodium methyl parahydroxybenzoate	2.000 mg
Sodium propyl parahydroxybenzoate	0.216 mg

White to off white suspension.

3. PACKAGE SIZE

1 Litre
2 Litres
5 Litres
10 Litres

4. TARGET SPECIES

Cattle, sheep, horses and other equines.

5. INDICATIONS FOR USE

Indications for use

For the treatment and control of gastro-intestinal roundworm infections in cattle, sheep and horses and of lungworm and tapeworm infections in cattle and sheep. This product is a broad spectrum anthelmintic for the control of all major benzimidazole-susceptible species of gastro-intestinal roundworms, including *Ostertagia* spp., *Teldorsagia* spp. and *Nematodirus* spp., and lungworms affecting cattle and sheep. It is effective against the adult and immature stages of gastro-intestinal roundworms and lungworms and also kills roundworm eggs. The product is also usually effective against Type II winter ostertagiasis in cattle and against *Moniezia* spp. of tapeworm in cattle and sheep. In horses and other equines, the product is effective against benzimidazole-susceptible large and small strongyles, ascarids, *Oxyuris* spp. and *Strongyloides* spp.

6. CONTRAINDICATIONS

Contraindications

Do not use in horses and other equines intended for human consumption.
Fenbendazole as a medicated liquid feed should not be used in the treatment of clinical infestations in cattle and sheep.

7. SPECIAL WARNINGS

Special warnings

Special warnings:
When administered by divided dosage in the form of liquid feed, the product may not be effective against *Strongyloides* spp. and *Trichuris* spp. in cattle and *Strongyloides* spp., *Dictyocaulus* spp. and *Bunostomum* spp. in sheep.

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 bodyweight, 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 anthelmintic, an anthelmintic belonging to another pharmacological class and having a different mode of action should be used.
Resistance to benzimidazoles (which include fenbendazole) has been reported in *Teladorsagia* spp., *Haemonchus* spp., *Cooperia* spp. and *Trichostrongylus* spp. in small ruminants in a number of countries, including the EU. Resistance to fenbendazole has been reported in cyathostomes in horses. Therefore, the use of this 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:
Intensive use or misuse of anthelmintics can give rise to resistance. To reduce this risk, dosing programmes should be discussed with your veterinary surgeon.

Special precautions to be taken by the person administering the veterinary medicinal product to animals:
Do not smoke, eat or drink when handling the veterinary medicinal product.
Avoid contact with the skin, eyes and mucous membranes. In case of accidental spillage onto the skin, eyes or mucous membranes, wash skin thoroughly with soap and water and rinse eyes and mucous membranes with plenty of water.
Personal protective equipment including impermeable rubber gloves should be worn when handling the veterinary medicinal product.
In case of accidental ingestion, seek medical advice immediately and show the package leaflet or the label to the physician.
Wash hands thoroughly with soap and water after use.

Pregnancy:
Can be used during pregnancy. Pregnant mares and young foals may also be safely treated with fenbendazole at the therapeutic dosage levels.

Interactions with other medicinal products and other forms of interaction:
None known.

Overdose:
Benzimidazoles have a high margin of safety. No specific overdose symptoms are known. No specific action is required.

Major incompatibilities:
None known.

8. ADVERSE EVENTS

Adverse events
Cattle, sheep, 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 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 or the local representative of the marketing authorisation holder using the contact details on this label, or via your national reporting system:
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
Oral use. Shake container before use.

Cattle and Horses: 1 ml per 13 kg bodyweight (7.5 mg fenbendazole per kg bodyweight) to be given orally.

	Practical Dosage Recommendations Cattle and horses above 400 kg should be given a further 5 ml for each additional 65 kg bodyweight)					
Weight of cattle / horses	65 kg	135 kg	200 kg	265 kg	335 kg	400 kg
Dose	5 ml	10 ml	15 ml	20 ml	25 ml	30 ml

Horses:
Recommended dosage 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.

Five day course: For the treatment and control of migrating and tissue larval stages of large strongyles, encysted mucosal 3rd and 4th stage small strongyle larvae and encysted inhibited 3rd stage small strongyle larvae in the mucosa administer 5 ml per 64 kg bodyweight daily for 5 days (7.5 mg fenbendazole per kg bodyweight daily for 5 days).

Single dose treatment: For the treatment and control of encysted mucosal stages of small strongyles administer 3 ml per 10 kg bodyweight (30 mg fenbendazole per kg bodyweight).
For the treatment and control of migrating and tissue stages of large strongyles administer 6 ml per 10 kg bodyweight (60 mg fenbendazole per kg bodyweight).
For the treatment of diarrhoea caused by *Strongyloides westeri* in two to three week old sucking foals administer 5 ml per 10 kg bodyweight (50 mg fenbendazole per kg bodyweight).

Sheep: 0.5 ml per 10 kg bodyweight (5 mg Fenbendazole per kg bodyweight) to be given orally.

	Practical Dosage Recommendations (Sheep above 60 kg should be given a further 0.5 ml for each additional 10 kg bodyweight)					
Weight of sheep	10 kg	20 kg	30 kg	40 kg	50 kg	60 kg
Dose	0.5 ml	1 ml	1.5 ml	2 ml	2.5 ml	3 ml

For administration to cattle and sheep a standard dosing gun or drenching equipment can be used.
For administration to horses, thoroughly mix the product with grain or concentrate

feed and give the full dosage as one administration.

Treatment should be repeated when reinfestation occurs.
Do not mix with other products.

10. ADVICE ON CORRECT ADMINISTRATION

Advice on correct administration
Underdosing could result in ineffective use and may favour resistance development.
To ensure a correct dosage, bodyweight should be determined as accurately as possible. Accuracy of the dosing device should be thoroughly checked.

11. WITHDRAWAL PERIODS

Withdrawal periods
Cattle:
Meat and offal: 12 days
Milk: 5 days
Sheep:
Meat and offal: 15 days
Milk: 7 days

Horses:
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.

12. SPECIAL STORAGE PRECAUTIONS

Special storage precautions
Keep out of the sight and reach of children.
Do not store above 25 °C.
Do not freeze.
Protect from frost.

Shelf life of the veterinary medicinal product as packaged for sale: 3 years.
Liquid feed containing the product will remain stable for up to 3 months.

13. SPECIAL PRECAUTIONS FOR DISPOSAL

Special precautions for disposal
Medicines should not be disposed of via wastewater.
This veterinary medicinal product should not enter water courses as fenbendazole may be dangerous for fish and other aquatic organisms.
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 06376/4076

Pack sizes
1, 2, 5 or 10 litre multidose containers.
Container: opaque white, high density polyethylene flat-bottle.
Closure: Tamper proof aluminium foil seal with polypropylene screw cap.

1 or 2.5 litre multidose containers.
Container: opaque white, high density polyethylene flexi-bottle with polypropylene screw cap.
Not all pack sizes may be marketed.

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

Manufacturer responsible for batch release:
Intervet Productions S.A.
Rue de Lyons
27460 Igoville
France

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

Distributor in Northern Ireland:
Intervet Ireland Ltd.
Magna Drive, Magna Business Park
Citywest Road, Dublin 24, Ireland

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

18. OTHER INFORMATION

Other information
Environmental properties
Fenbendazole is toxic to fish and other aquatic organisms.

POM-VPS Veterinary medicinal product subject to prescription.

Class of anthelmintic: 1-BZ

19. THE WORDS "FOR ANIMAL TREATMENT ONLY"

For animal treatment only.

20. EXPIRY DATE

Exp. {mm/yyyy}

21. BATCH NUMBER

Lot {number}

MINIMUM PARTICULARS TO APPEAR ON SMALL IMMEDIATE PACKAGING UNITS -Flexi-bottle; Front label 1 litre and 2.5 Litre

1. NAME OF THE VETERINARY MEDICINAL PRODUCT

Panacur 10% oral suspension

2. QUANTITATIVE PARTICULARS OF THE ACTIVE SUBSTANCES

Each ml contains:
Fenbendazole 100 mg

Cattle, sheep and horse wormer.

1 Litre
2.5 Litres

1 Litre:
1000 doses for 20 kg lambs
400 doses for 50 kg ewes
66 doses for 200 kg cattle
22 doses for 600 kg horses

2.5 Litres:
2500 doses for 20 kg lambs
1000 doses for 50 kg ewes
166 doses for 200 kg cattle
55 doses for 600 kg horses

3. BATCH NUMBER

Lot {number}

4. EXPIRY DATE

Exp. {mm/yyyy}

PARTICULARS TO APPEAR ON THE IMMEDIATE (OUTER) PACKAGE – 1 Litre and 2.5 Litre Flexi-bottle – BACK LABEL – (Base label and front cover of concertina label)

1. NAME OF THE VETERINARY MEDICINAL PRODUCT

Panacur 10% oral suspension

2. STATEMENT OF ACTIVE SUBSTANCES

Each ml contains:
Fenbendazole 100 mg

3. PACKAGE SIZE

1 Litre
2.5 Litres

4. TARGET SPECIES

Cattle, sheep, horses and other equines.

5. INDICATIONS

For the treatment and control of gastro-intestinal roundworm infections in cattle, sheep and horses and of lungworm and tapeworm infections in cattle and sheep.

6. ROUTES OF ADMINISTRATION

Oral use.

7. WITHDRAWAL PERIODS

Withdrawal periods:
Cattle:
Meat and offal: 12 days
Milk: 5 days

Sheep:
Meat and offal: 15 days
Milk: 7 days

Do not use in horses and other equines intended for human consumption.

8. EXPIRY DATE

Exp. {mm/yyyy}
Liquid feed containing the product will remain stable for up to 3 months.

9. SPECIAL STORAGE PRECAUTIONS

Do not store above 25 °C.
Do not freeze.
Protect from frost.

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

MA Holder:
Intervet International B.V.

Distributor in Northern Ireland:
Intervet Ireland Ltd.

14. MARKETING AUTHORISATION NUMBERS

Vm 06376/4076

15. BATCH NUMBER

Lot {number}

Class of anthelmintic: (1-BZ)

PARTICULARS TO APPEAR ON THE PACKAGE LEAFLET:

PACKAGE LEAFLET

1. Name of the veterinary medicinal product

Panacur 10% oral suspension

2. Composition

Each ml contains:
Active substance:

Fenbendazole	100 mg

Excipients:

Benzyl alcohol	4.835 mg
Sodium methyl parahydroxybenzoate	2.000 mg
Sodium propyl parahydroxybenzoate	0.216 mg

White to off white suspension.

3. Target species

Cattle, sheep, horses and other equines.

4. Indications for use

For the treatment and control of gastro-intestinal roundworm infections in cattle, sheep and horses and of lungworm and tapeworm infections in cattle and sheep. This product is a broad spectrum anthelmintic for the control of all major benzimidazole-susceptible species of gastro-intestinal roundworms, including *Ostertagia* spp., *Teladorsagia* spp. and *Nematodirus* spp. and lungworms affecting cattle and sheep. It is effective against the adult and immature stages of gastro-intestinal roundworms and lungworms and also kills roundworm eggs. The product is also usually effective against Type II winter ostertagiasis in cattle and against *Moniezia* spp. of tapeworm in cattle and sheep. In horses and other equines, the product is effective against benzimidazole-susceptible large and small strongyles, ascarids, *Oxyuris* spp. and *Strongyloides* spp.

5. Contraindications

Do not use in horses and other equines intended for human consumption.
Fenbendazole as a medicated liquid feed should not be used in the treatment of clinical infestations in cattle and sheep.

6. Special warnings

Special warnings:
When administered by divided dosage in the form of liquid feed, the product may not be effective against *Strongyloides* spp. and *Trichuris* spp. in cattle and *Strongyloides* spp., *Dictyocaulus* spp. and *Bunostomum* spp. in sheep.

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 bodyweight, 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 anthelmintic, an anthelmintic belonging to another pharmacological class and having a different mode of action should be used.
Resistance to benzimidazoles (which include fenbendazole) has been reported in *Teladorsagia* spp., *Haemonchus* spp., *Cooperia* spp. and *Trichostrongylus* spp. in small ruminants in a number of countries, including the EU. Resistance to fenbendazole has been reported in cyathostomes in horses. Therefore, the use of this 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:
Intensive use or misuse of anthelmintics can give rise to resistance. To reduce this risk, dosing programmes should be discussed with your veterinary surgeon.

Special precautions to be taken by the person administering the veterinary medicinal product to animals:
Do not smoke, eat or drink when handling the veterinary medicinal product.
Avoid contact with the skin, eyes and mucous membranes. In case of accidental spillage onto the skin, eyes or mucous membranes, wash skin thoroughly with soap and water and rinse eyes and mucous membranes with plenty of water.
Personal protective equipment including impermeable rubber gloves should be worn when handling the veterinary medicinal product.
In case of accidental ingestion, seek medical advice immediately and show the package leaflet or the label to the physician.
Wash hands thoroughly with soap and water after use.

Pregnancy:
Can be used during pregnancy. Pregnant mares and young foals may also be safely treated with fenbendazole at the therapeutic dosage levels.

Interaction with other medicinal products and other forms of interaction:
None known.

Overdose:
Benzimidazoles have a high margin of safety. No specific overdose symptoms are known. No specific action is required.

Major incompatibilities:
None known.

7. Adverse events

Cattle, sheep, 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: 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. Shake container before use.
Cattle and Horses: 1 ml per 13 kg bodyweight (7.5 mg fenbendazole per kg bodyweight) to be given orally (see table).

	Practical Dosage Recommendations (Cattle and horses above 400 kg should be given a further 5 ml for each additional 65 kg bodyweight)					
Weight of cattle / horses	65 kg	135 kg	200 kg	265 kg	335 kg	400 kg
Dose	5 ml	10 ml	15 ml	20 ml	25 ml	30 ml

Horses:
Recommended dosage 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.

Five day course: For the treatment and control of migrating and tissue larval stages of large strongyles, encysted mucosal 3rd and 4th stage small strongyle larvae and encysted inhibited 3rd stage small strongyle larvae in the mucosa administer 5 ml per 64 kg bodyweight daily for 5 days (7.5 mg fenbendazole per kg bodyweight daily for 5 days).

Single dose treatment: For the treatment and control of encysted mucosal stages of small strongyles administer 3 ml per 10 kg bodyweight (30 mg fenbendazole per kg bodyweight).
For the treatment and control of migrating and tissue stages of large strongyles administer 6 ml per 10 kg bodyweight (60 mg fenbendazole per kg bodyweight).
For the treatment of diarrhoea caused by *Strongyloides westeri* in two to three week old sucking foals administer 5 ml per 10 kg bodyweight (50 mg fenbendazole per kg bodyweight).

Sheep: 0.5 ml per 10 kg bodyweight (5 mg fenbendazole per kg bodyweight) to be given orally (see table).

	Practical Dosage Recommendations (Sheep above 60 kg should be given a further 0.5 ml for each additional 10 kg bodyweight)					
Weight of sheep	10 kg	20 kg	30 kg	40 kg	50 kg	60 kg
Dose	0.5 ml	1 ml	1.5 ml	2 ml	2.5 ml	3 ml

For administration to cattle and sheep a standard dosing gun or drenching equipment can be used.
For administration to horses, thoroughly mix the product with grain or concentrate feed and give the full dosage as one administration.
Treatment should be repeated when reinfestation occurs.
Do not mix with other products.

9. Advice on correct administration

Underdosing could result in ineffective use and may favour resistance development.
To ensure a correct dosage, bodyweight should be determined as accurately as possible. Accuracy of the dosing device should be thoroughly checked.

10. Withdrawal periods

Cattle:
Meat and offal: 12 days
Milk: 5 days
Sheep:
Meat and offal: 15 days
Milk: 7 days

Horses:
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.
Do not store above 25 °C.
Do not freeze.
Protect from frost.

Shelf life of the veterinary medicinal product as packaged for sale: 3 years.
Liquid feed containing the product will remain stable for up to 3 months.

12. Special precautions for disposal

Medicines should not be disposed of via wastewater.
This veterinary medicinal product should not enter water courses as fenbendazole may be dangerous for fish and other aquatic organisms.
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 06376/4076

Pack sizes:
1, 2, 5 or 10 litre multidose containers.
Container: opaque white, high density polyethylene flat-bottle. Closure: Tamper proof aluminium foil seal with polypropylene screw cap.

1 or 2.5 litre multidose containers.
Container: opaque white, high density polyethylene flexi-bottle with polypropylene screw cap.
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 Productions S.A.
Rue de Lyons
27460 Igoville
France

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

Distributor in Northern Ireland:
Intervet Ireland Ltd.
Magna Drive, Magna Business Park
Citywest Road, Dublin 24, Ireland

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

17. Other information

Environmental properties
Fenbendazole is toxic to fish and other aquatic organisms.

POM-VPS Veterinary medicinal product subject to prescription.

Class of anthelmintic: 1-BZ

Gavin Hall
Approved: 25 July 2025