

PARTICULARS TO APPEAR ON THE IMMEDIATE PACKAGE {Bottle labels}

1. NAME OF THE VETERINARY MEDICINAL PRODUCT

Zermex 0.1% w/v Oral Solution for Sheep

Moxidectin

2. STATEMENT OF ACTIVE SUBSTANCES

Moxidectin 1 mg/ml

3. PHARMACEUTICAL FORM

Oral Solution for Sheep

4. PACKAGE SIZE

1 L

2.5 L

5 L

5. TARGET SPECIES

Sheep

6. INDICATION(S)

For the treatment and control of:

Adult and immature roundworms including: *Nematodirus*, plus inhibited larvae of *Ostertagia*, *Haemonchus* and *Trichostrongylus*

Benzimidazole resistant strains of:

Ostertagia, *Haemonchus*, *Trichostrongylus* and *Cooperia*

ZERMEX ORAL SOLUTION has a persistent effect in preventing re-infestation by *Haemonchus contortus* and *Teladorsagia circumcincta* for 5 weeks and *Oesophagostomum columbianum* for 4 weeks after treatment.

7. METHOD AND ROUTE(S) OF ADMINISTRATION

ZERMEX ORAL SOLUTION is ready-to-use. Administer 1 ml per 5 kg liveweight using the ZERMEX Drench Gun or any standard drenching equipment. Check dose rates and equipment before drenching commences.

READ PACKAGE LEAFLET BEFORE USE

8. WITHDRAWAL PERIOD

Meat and offal: 14 days.

Milk: 5 days.

9. SPECIAL WARNING(S), IF NECESSARY

ZERMEX ORAL SOLUTION has been shown to be safe for use in pregnant, lactating and breeding animals.

Operator warnings:

Wear protective impermeable gloves during use. Avoid direct contact with skin and eyes. In case of contact with skin and eyes wash affected area with clean water. Seek medical advice if irritation persists. Do not smoke, drink or eat while handling the veterinary medicinal product. Wash hands after use.

Environmental risks have been identified for this product and special precautions apply. Read the package leaflet before use.

10. EXPIRY DATE

Expiry date:{month/year}

Once opened use within 6 months.

Once opened, used by:

11. SPECIAL STORAGE CONDITIONS

Do not store above 25°C. Protect from light.

Keep container in the outer carton.

12. SPECIFIC PRECAUTIONS FOR THE DISPOSAL OF UNUSED PRODUCTS OR WASTE MATERIALS, IF ANY

Any unused veterinary medicinal product or waste material derived from such veterinary medicinal products should be disposed of in accordance with local requirements. Do not contaminate watercourses with the product. DANGEROUS to fish and aquatic life.

13. THE WORDS “FOR ANIMAL TREATMENT ONLY” AND CONDITIONS OR RESTRICTIONS REGARDING SUPPLY AND USE IF APPLICABLE

POM-VPS

To be supplied only on veterinary prescription.

FOR ANIMAL TREATMENT ONLY

3-ML

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

KEEP OUT OF THE SIGHT AND REACH OF CHILDREN

15. NAME AND ADDRESS OF THE MARKETING AUTHORISATION HOLDER

Zoetis Belgium S.A.
2nd Floor, Building 10
Cherrywood Business Park
Loughlinstown
Dublin 18
D18 T3Y1
Ireland

Distributor:
Downland Marketing Limited
Main Mill
Warwick Mill Business Centre
Warwick Bridge
Carlisle
Cumbria
CA4 8RR

16. MARKETING AUTHORISATION NUMBER

Vm 60021/3029

17. MANUFACTURER’S BATCH NUMBER

Batch No:

PARTICULARS TO APPEAR ON THE OUTER PACKAGE {Carton}

1. NAME OF THE VETERINARY MEDICINAL PRODUCT

Zermex 0.1% w/v Oral Solution for Sheep

Moxidectin

2. STATEMENT OF ACTIVE SUBSTANCES

Moxidectin 1 mg/ml

3. PHARMACEUTICAL FORM

Oral Solution

4. PACKAGE SIZE

1 L

2.5 L

5 L

5. TARGET SPECIES

Sheep

6. INDICATION(S)

For the treatment and control of:

Adult and immature roundworms including: *Nematodirus*, plus inhibited larvae of *Ostertagia*, *Haemonchus* and *Trichostrongylus*

Benzimidazole resistant strains of:

Ostertagia, *Haemonchus*, *Trichostrongylus* and *Cooperia*

ZERMEX ORAL SOLUTION is effective against the following parasites of sheep.

Effective against adult and immature (L4) GI worms

- *Teladorsagia circumcincta* (including inhibited larvae)
- *Ostertagia trifurcate*
- *Haemonchus contortus* (including inhibited larvae)
- *Trichostrongylus axei* (including inhibited larvae)
- *Trichostrongylus colubriformis*
- *Nematodirus spathiger*
- *Nematodirus filicollis* (adults only)
- *Strongyloides papillosus* (larval stages only)
- *Cooperia curticei* (adults only)
- *Cooperia oncophora*
- *Oesophagostomum columbianum*

- *Trichostrongylus vitrinus*
- *Nematodirus battus*
- *Oesophagostomum venulosum* (adults only)
- *Chabertia ovina*
- *Trichuris ovis* (adults only)

Effective against adult lungworm

- *Dictyocaulus filaria*

ZERMEX ORAL SOLUTION has a persistent effect in preventing re-infection by *Haemonchus contortus* and *Ostertagia circumcincta* for 5 weeks and *Oesophagostomum columbianum* for 4 weeks after treatment.

7. METHOD AND ROUTE(S) OF ADMINISTRATION

ZERMEX ORAL SOLUTION is ready-to-use. Administer 1 ml per 5 kg liveweight using the ZERMEX Drench Gun or any standard drenching equipment. Check dose rates and equipment before drenching commences.

1 ml/5 kg liveweight to give a dose of 0.2 mg moxidectin/kg liveweight.

Liveweight (kg)	Dose volume (ml)	Doses per Pack (1L)	Doses per Pack (2.5L)	Doses per Pack (5L)
10	2	500	1250	2500
11-15	3	333	833	1666
16-20	4	250	625	1250
21-25	5	200	500	1000
26-30	6	166	416	833
31-35	7	142	357	714
36-40	8	125	312	625
41-45	9	111	277	555
46-50	10	100	250	500
51-55	11	90	227	454
56-60	12	83	208	416
61-65	13	76	192	384
66-70	14	71	178	357
71-75	15	66	166	333

For animals over 75 kg liveweight, dose at 1 ml per 5 kg.

8. WITHDRAWAL PERIOD

Meat and offal: 14 days.

Milk: 5 days.

9. SPECIAL WARNING(S), IF NECESSARY

ZERMEX ORAL SOLUTION has been shown to be safe for use in pregnant, lactating and breeding animals.

Do not exceed stated dose. Do not mix with other veterinary medicinal products before administration.

Operator warnings:

Wear protective impermeable gloves during use. Avoid direct contact with skin and eyes. In case of contact with skin and eyes wash affected area with clean water. Seek medical advice if irritation persists. Do not smoke, drink or eat while handling the veterinary medicinal product. Wash hands after use.

Environmental risks have been identified for this product and special precautions apply. Read the package leaflet before use.

10. EXPIRY DATE

Expiry date: {month/year}

Once opened used within 6 months.

Once opened, use by:

11. SPECIAL STORAGE CONDITIONS

Do not store above 25°C. Protect from light.

Keep container in the outer carton.

12. SPECIFIC PRECAUTIONS FOR THE DISPOSAL OF UNUSED PRODUCTS OR WASTE MATERIALS, IF ANY

Any unused veterinary medicinal product or waste material derived from such veterinary medicinal products should be disposed of in accordance with local requirements. Do not contaminate watercourses with the product. DANGEROUS to fish and aquatic life.

13. THE WORDS “FOR ANIMAL TREATMENT ONLY” AND CONDITIONS OR RESTRICTIONS REGARDING SUPPLY AND USE IF APPLICABLE

POM-VPS

To be supplied only on veterinary prescription

FOR ANIMAL TREATMENT ONLY

3-ML

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

KEEP OUT OF THE SIGHT AND REACH OF CHILDREN

15. NAME AND ADDRESS OF THE MARKETING AUTHORISATION HOLDER

Zoetis Belgium S.A.
2nd Floor, Building 10
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Cumbria
CA4 8RR

16. MARKETING AUTHORISATION NUMBER

Vm 60021/3029

17. MANUFACTURER'S BATCH NUMBER

Batch No:

PACKAGE LEAFLET FOR: Zermex 0.1% w/v Oral Solution for Sheep

1. NAME AND ADDRESS OF THE MARKETING AUTHORISATION HOLDER AND OF THE MANUFACTURING AUTHORISATION HOLDER RESPONSIBLE FOR BATCH RELEASE, IF DIFFERENT

MA Holder:

Zoetis Belgium S.A.
2nd Floor, Building 10
Cherrywood Business Park
Loughlinstown
Dublin 18
D18 T3Y1
Ireland

Manufacturing site responsible for batch release:

Zoetis Manufacturing & Research Spain S.L.
Carretera Camprodon s/n – La Riba
17813 – Vall de Bianya
Girona
Spain

2. NAME OF THE VETERINARY MEDICINAL PRODUCT

ZERMEX 0.1% W/V ORAL SOLUTION FOR SHEEP

Moxidectin

3. STATEMENT OF THE ACTIVE SUBSTANCE (S) AND OTHER INGREDIENTS

Each ml contains:

Active substance:

Moxidectin	1 mg
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Excipients:

Benzyl Alcohol	40 mg
Butylated Hydroxytoluene	2.5 mg

A pale, yellow solution

4. INDICATION(S)

ZERMEX ORAL SOLUTION FOR SHEEP is a second generation macrocyclic lactone parasiticide of the milbemycin family which is effective against internal parasites of sheep. ZERMEX belongs to the avermectin/milbemycin 3-ML class of anthelmintics.

ZERMEX ORAL SOLUTION treats and controls the following parasites of sheep.

Effective against adult and immature (L4) GI worms

- *Teladorsagia circumcincta* (including inhibited larvae)
- *Ostertagia trifurcata*
- *Haemonchus contortus* (including inhibited larvae)
- *Trichostrongylus axei* (including inhibited larvae)
- *Trichostrongylus colubriformis*
- *Trichostrongylus vitrinus*
- *Nematodirus battus*
- *Nematodirus spathiger*
- *Nematodirus filicollis* (adults only)
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- *Cooperia curticei* (adults only)
- *Cooperia oncophora*
- *Oesophagostomum columbianum*
- *Oesophagostomum venulosum* (adults only)
- *Chabertia ovina*
- *Trichuris ovis* (adults only)

Effective against adult lungworm

- *Dictyocaulus filaria*

ZERMEX ORAL SOLUTION has persistent effect in preventing re-infection by *Haemonchus contortus* and *Ostertagia circumcincta* for 5 weeks and *Oesophagostomum columbianum* for 4 weeks after treatment.

5. CONTRAINDICATIONS

Do not mix with other products.

6. ADVERSE REACTIONS None known.

7. TARGET SPECIES

Sheep

8. DOSAGE FOR EACH SPECIES, ROUTE(S) AND METHOD OF ADMINISTRATION

Administer 1 ml per 5 kg liveweight using the ZERMEX Drench Gun or any standard drenching equipment.

1 ml/5 kg liveweight to give a dose of 0.2 mg moxidectin/kg liveweight.

Liveweight (kg)	Dose volume (ml)	Doses per Pack (1L)	Doses per Pack (2.5L)	Doses per Pack (5L)
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56-60	12	83	208	416
61-65	13	76	192	384
66-70	14	71	178	357
71-75	15	66	166	333

For animals over 75 kg liveweight, dose at 1 ml per 5 kg.

9. ADVICE ON CORRECT ADMINISTRATION

ZERMEX ORAL DRENCH is ready-to-use.

Underdosing could result in ineffective use and may favour resistance development.

To ensure administration of a correct dose, bodyweight should be determined as accurately as possible. If animals are to be treated collectively, reasonably homogenous groups should be set up, and all animals of a group should be dosed at the rate corresponding to the heaviest one. Accuracy of the dosing device should be thoroughly checked.

Do not mix with other veterinary medicinal products before administration. Avoid the introduction of contamination during use.

10. WITHDRAWAL PERIOD(S)

Meat and offal: 14 days

Milk: 5 days

11. SPECIAL STORAGE PRECAUTIONS

Keep out of the sight and reach of children.

Do not store above 25°C. Protect from light. Keep container in outer carton.

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

Shelf life after first opening the container: 6 months

When the container is broached/opened for the first time, using the in-use shelf-life which is specified on this package leaflet, the date on which any product remaining in the container should be discarded should be determined. This discard date should be written in the space provided on the label.

12. SPECIAL WARNING(S)

Unnecessary use of antiparasitics or use deviating from the instructions given in the SPC may increase the resistance selection pressure and lead to reduced efficacy. The decision to use the product should be based on confirmation of the parasitic species and burden, or of the risk of infection based on its epidemiological features, for each flock.

Repeated use for an extended period, particularly when using the same class of substances, increases the risk of resistance development. Within a flock, maintenance of susceptible refugia is essential to reduce that risk. Systematically applied interval-based treatment and treatment of a whole flock should be avoided. Instead, if feasible, only selected individual animals or subgroups should be treated (targeted selective treatment). This should be combined with appropriate husbandry and pasture management measures. Guidance for each specific flock should be sought from the responsible veterinarian.

Multiple resistance of *Teladorsagia circumcincta* to moxidectin, levamisole, benzimidazole and ivermectin was reported throughout Europe. Moxidectin-resistant *Haemonchus contortus* and *Trichostrongylus colubriformis* were also described.. Therefore the use of this product should take into account local information about susceptibility of the target parasites, where available. Additionally, use should be based on local history of treatments and recommendations on how to use the product under sustainable conditions to limit further selection for resistance to anthelmintics. These precautions are especially important when moxidectin is being used to control resistant strains.

Clinical trials have shown that ZERMEX ORAL SOLUTION is effective against certain benzimidazole (white drench) resistant strains of *Haemonchus*, *Ostertagia*, *Trichostrongylus* and *Cooperia*.

It is recommended to further investigate cases of suspected resistance, using an appropriate diagnostic method (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. Confirmed resistance should be reported to the marketing authorisation holder or to the competent authorities.

Symptoms of overdose generally do not occur at less than 5 times the recommended dose. They are manifested as transient salivation, depression, drowsiness and ataxia 8 to 12 hours post-treatment. Treatment is not generally necessary and recovery is generally complete within 24 to 48 hours. There is no specific antidote.

User warnings:

Wear protective impermeable gloves during use. Avoid direct contact with skin and eyes. In case of contact with skin and eyes wash affected area with clean water. Seek medical advice if irritation persists. Do not smoke, drink or eat while handling the veterinary medicinal product. Wash hands after use.

Other precautions regarding impact on the environment:

Moxidectin fulfils the criteria for a (very) persistent, bioaccumulative and toxic (PBT) substance; therefore, exposure of the environment to moxidectin must be limited to the extent possible. Treatments should be administered only when necessary and should be based on faecal egg counts or evaluation of the risk of infestation at the animal and/or flock level.

Like other macrocyclic lactones, moxidectin has the potential to adversely affect non-target organisms:

- Faeces containing moxidectin excreted onto pasture by treated animals may temporarily reduce the abundance of dung feeding organisms. Following treatment of sheep with the product, levels of moxidectin that are potentially toxic to dung fly species may be excreted over a period of 4 days and may decrease dung fly abundance during that period. It has been established in laboratory tests that moxidectin may temporarily affect dung beetle reproduction; however, studies with incurred residues indicate no long-term effects. Nevertheless, in case of repeated treatments with moxidectin (as with products of the same anthelmintic class) it is advisable not to treat animals every time on the same pasture to allow dung fauna populations to recover.
- Moxidectin is inherently toxic to aquatic organisms including fish. The product should be used only according to the label instructions. Based on the excretion profile of moxidectin when administered as the oral formulation to sheep, treated animals should not have access to watercourses during the first 3 days after treatment.

Incompatibilities

In the absence of compatibility studies, this veterinary medicinal product must not be mixed with other veterinary medicinal products.

13. SPECIAL PRECAUTIONS FOR THE DISPOSAL OF UNUSED PRODUCT OR WASTE MATERIALS, IF ANY

Any unused veterinary medicinal product or waste material derived from such veterinary medicinal products should be disposed of in accordance with local requirements. Do not contaminate watercourses with the product. DANGEROUS to fish and aquatic life.

Ask your veterinary surgeon or pharmacist how to dispose of medicines no longer required. These measures should help to protect the environment.

14. 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.

15. OTHER INFORMATION

For animal treatment only.

POM-VPS

To be supplied only on veterinary prescription.

Vm 60021/3029

3-ML

Distributed in the UK by:
Downland Marketing Limited
Main Mill
Warwick Mill Business Centre
Warwick Bridge
Carlisle
Cumbria
CA4 8RR

ZERMEX ORAL SOLUTION has been shown to be safe for use in pregnant, lactating and breeding animals.

ZERMEX ORAL SOLUTION is packaged in 1, 2.5 and 5L HDPE bottles with PP screw cap closures. Not all pack sizes may be marketed.

Gavin Hall

Approved: 14 January 2025