

A. SYRINGE LABEL

PARTICULARS TO APPEAR ON THE IMMEDIATE PACKAGE

Syringe

1. NAME OF THE VETERINARY MEDICINAL PRODUCT

Animec 18.7 mg/g Oral Paste for Horses

Ivermectin

2. STATEMENT OF ACTIVE AND OTHER SUBSTANCES

Ivermectin 18.7 mg/g

3. PHARMACEUTICAL FORM

Oral Paste

4. PACKAGE SIZE

6.42 g or 7.49 g

5. TARGET SPECIES

Horses

6. INDICATION(S)

7. METHOD AND ROUTE(S) OF ADMINISTRATION

Antiparasitic treatment for Horses.

Read the package leaflet or outer carton before use.

8. WITHDRAWAL PERIOD

Meat and offal: 34 days
Do not use in mares producing milk for human consumption

9. SPECIAL WARNING(S), IF NECESSARY

10. EXPIRY DATE

MM/YYYY

The product is for single use. After use the syringe should be discarded

11. SPECIAL STORAGE CONDITIONS

Do not store above 25 °C

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

13. THE WORDS “FOR ANIMAL TREATMENT ONLY” AND CONDITIONS OR RESTRICTIONS REGARDING SUPPLY AND USE, if applicable

For animal treatment only

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

Keep out of the reach and sight of children

15. NAME AND ADDRESS OF THE MARKETING AUTHORISATION HOLDER

ACME DRUGS s.r.l.
Via Portella della Ginestra 9
42025 Cavriago
Italy

16. MARKETING AUTHORISATION NUMBER

United Kingdom

Veterinary Medicinal Product authorised for use in UK and IE.

UK ONLY	
POM-VPS	To be supplied only on a veterinary prescription
	Vm 61301/5000

17. MANUFACTURER'S BATCH NUMBER

B. BOX LABEL

PARTICULARS TO APPEAR ON THE OUTER PACKAGE

Box

1. NAME OF THE VETERINARY MEDICINAL PRODUCT

Animec 18.7 mg/g Oral Paste for Horses
Ivermectin

2. STATEMENT OF ACTIVE AND OTHER SUBSTANCES

1 Syringe containing 7.49 g (6.42 g) paste which contains 18.7 mg/g ivermectin

3. PHARMACEUTICAL FORM

Oral Paste

4. PACKAGE SIZE

6.42 g or 7.49g

5. TARGET SPECIES

Horses

6. INDICATION(S)

Antiparasitic treatment for Horses.

7. METHOD AND ROUTE(S) OF ADMINISTRATION

One syringe will treat upto a 700kg horse (only for 7.49 g syringe)
Read the package leaflet before use.

8. WITHDRAWAL PERIOD

Meat and offal: 34 days
Do not use in mares producing milk for human consumption.

9. SPECIAL WARNING(S), IF NECESSARY

Read the package leaflet before use for full instructions and user warnings.

10. EXPIRY DATE

MM/YYYY

The product is for single use. After use the syringe should be discarded.

11. SPECIAL STORAGE CONDITIONS

Do not store above 25°C

12. SPECIAL PRECAUTIONS FOR THE DISPOSAL OF UNUSED PRODUCTS

13. THE WORDS "KEEP OUT OF THE REACH AND SIGHT OF CHILDREN" AND CONDITIONS OR RESTRICTIONS REGARDING SUPPLY AND USE, if applicable

For animal treatment only

14. THE WORDS "KEEP OUT OF THE REACH AND SIGHT OF CHILDREN"

Keep out of the reach and sight of children

15. NAME AND ADDRESS OF THE MARKETING AUTHORISATION HOLDER

ACME DRUGS s.r.l.
Via Portella della Ginestra 9
42025 Cavriago
Italy

16. MARKETING AUTHORISATION NUMBER

United Kingdom
Veterinary Medicinal Product authorised for use in UK and IE.

UK ONLY
To be supplied only on a
veterinary prescription
Vm 61301/5000

POM-VPS

17. MANUFACTURER'S BATCH NUMBER

C. PACKAGE LEAFLET

PACKAGE LEAFLET

Animec 18.7 mg/g Oral Paste for Horses

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

Marketing authorisation holder

ACME DRUGS s.r.l.
Via Portella della Ginestra 9
42025 Cavriago
Italy

Manufacturer for the batch release:

ACME Drugs S.r.l.
Via Portella della Ginestra, 9, Zona Industriale Corte Tegge
42025 CAVRIAGO (RE)
Italy

2. NAME OF THE VETERINARY MEDICINAL PRODUCT

Animec 18.7 mg/g Oral Paste for Horses

Ivermectin

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

Ivermectin 18.7 mg/g
A white homogeneous paste

4. INDICATION(S)

Treatment of nematode or arthropod infection due to:

Large strongyles:

Strongylus vulgaris (adults and L₄ stage larvae [arterial])
Strongylus edentatus (adults and L₄ stage larvae [tissue])
Strongylus equinus (adults)

Small strongyles (including benzimidazole resistant strains):

Cyathostomum spp (adults and luminal L₄ stage larvae)
Cylicocyclus spp. (adults and luminal L₄ stage larvae)
Cylicodontophorus spp. (adults and luminal L₄ stage larvae)
Cylicostephanus spp. (adults and luminal L₄ stage larvae)
Gyalocephalus spp. (adults and luminal L₄ stage larvae)

Ascarids:
Parascaris equorum (luminal L₅ larvae and adults)

Pinworms:
Oxyuris equi (L₄ stage larvae and adults)

Neck threadworms:
Onchocerca spp (microfilariae)

Stomach bots:
Gasterophilus spp (oral and gastric stages)

5. CONTRAINDICATIONS

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

Do not use in dogs or cats as severe adverse reactions may occur.

6. ADVERSE REACTIONS

The effects of GABA agonists are increased by ivermectin

If you notice any serious effects or other effects not mentioned in this leaflet, please inform your veterinary surgeon

7. TARGET SPECIES

Horses

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

Dosage:

One syringe division of paste per 100 kg body weight (based on a recommended dosage of 200 microgram ivermectin per kg body weight).. The syringe containing 6.42 g of paste delivers sufficient paste to treat 600 kg of bodyweight at the recommended dose rate.

The syringe containing 7.49 g of paste delivers sufficient paste to treat 700 kg of bodyweight at the recommended dose rate.

Administration:

The paste is given by oral route.

9. ADVICE ON CORRECT ADMINISTRATION

To ensure administration of a correct dose, body weight should be determined as accurately as possible. The animal's mouth should be free from food to ensure

swallowing. Turn the screw gauge on the syringe plunger to the body weight of the horse. The tip of the syringe barrel should be inserted at the interdental space (the gap between the front and back teeth) and the paste deposited on the base of the tongue. Advance the plunger as far as it will go, depositing the medication on the

base of the tongue. Immediately elevate the horse's head for a few seconds to ensure swallowing.

10. WITHDRAWAL PERIOD

Meat and offal: 34 days

Do not use in mares producing milk for human consumption.

11. SPECIAL STORAGE PRECAUTIONS

Keep out of the reach and sight of children.

Do not store above 25°C

The product is for single use. After use, the syringe should be discarded.

Do not use after expiry date which is stated on the label/carton

12. SPECIAL WARNING(S)

Special precautions for use in animals

Some horses with heavy infections of *Onchocerca* spp. microfilariae have experienced oedema and pruritus following treatment. Such reactions are assumed to be the result of death of large numbers of microfilariae. These signs resolve within a few days but symptomatic treatment may be advisable

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 or misadministration of the product.

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.

Veterinary advice should be given on appropriate dosing programs and stock management to achieve adequate parasite control and reduce the likelihood of anthelmintic resistance developing. In the event that a product is suspected of being ineffective, the animal owner is advised to seek veterinary advice.

Resistance to ivermectin has been reported in *Parascaris equorum*. Therefore, the use of this product should be based on local farm epidemiological information about

susceptibility of nematodes and recommendations on how to limit further selection for resistance to anthelmintics

Since ivermectin is highly bound to plasma proteins, special care should be taken in cases of sick animals or in nutritional conditions associated with low plasma protein levels.

Dogs and cats should not be allowed to ingest spilled gel or have access to used packaging due to the potential for adverse effects related to ivermectin toxicity

The product has been formulated for use in horses only. Cats, dogs, especially Collies, Old English Sheepdogs and related breeds or crosses, and also turtles and tortoises may be adversely affected by the concentration of ivermectin in this product if they are allowed to ingest spilled paste or have access to used syringes.

The product can be administered to mares at any stages of pregnancy or lactation.

Mild transitory signs (slowed pupillary light response and depression) have been seen at a higher dose of 1.8 mg/kg (9 times the recommended dose level). Other signs seen at higher doses include mydriasis (dilation of the pupil), ataxia (incoordination), tremors (shaking), stupor (dull), coma and death. The less severe signs have been transitory.

Although no antidote has been identified, symptomatic therapy may be beneficial.

User warnings

Do not smoke, eat or drink while handling the product.

Wash hands after use.

This product may cause skin and eye irritation. Therefore, the user should avoid contact of the product with the skin and the eyes. In case of contact, rinse immediately with plenty of water.

In the case of accidental ingestion or eye irritation after contact seek medical advice immediately and show the package insert or the label to the physician.

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

EXTREMELY DANGEROUS TO FISH AND AQUATIC LIFE. Do not contaminate surface waters or ditches with product or used container. Any unused veterinary medicinal product or waste materials derived from such veterinary medicinal products should be disposed of in accordance with local requirements.

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

Veterinary Medicinal Product authorised for use in UK and IE.

Pack size:

Box containing 1 syringe of 6,42 g

Box containing 1 syringe of 7,49 g

Box containing 50 syringes of 7,49 g

Not all pack sizes may be marketed

UK ONLY

POM-
VPS

—
To be supplied only on a veterinary prescription

Vm 61301/5000

Dispose of any unused product and empty containers in accordance with guidance from your local waste regulation authority

B. BOX LABEL
Box of 50 syringes

PARTICULARS TO APPEAR ON THE OUTER PACKAGE

Box of 50 syringes

1. NAME OF THE VETERINARY MEDICINAL PRODUCT

Animec 18.7 mg/g Oral Paste for Horses

Ivermectin

2. STATEMENT OF ACTIVE AND OTHER SUBSTANCES

Ivermectin 18.7 mg/g

A white homogeneous paste

3. PHARMACEUTICAL FORM

Oral Paste

4. PACKAGE SIZE

Each syringe containing 7.49 g paste which contains 18.7 mg/g ivermectin 50 syringes per box

5. TARGET SPECIES

Horses

6. INDICATION(S)

Treatment of nematode or arthropod infection due to:

Large strongyles:

Strongylus vulgaris (adults and L₄ stage larvae [arterial])

Strongylus edentatus (adults and L₄ stage larvae [tissue])

Strongylus equinus (adults)

Small strongyles (including benzimidazole resistant strains):

Cyathostomum spp (adults and luminal L₄ stage larvae)

Cylicocyclus spp. (adults and luminal L₄ stage larvae)

Cylicodontophorus spp. (adults and luminal L₄ stage larvae)

Cylicostephanus spp. (adults and luminal L₄ stage larvae)
Gyalocephalus spp. (adults and luminal L₄ stage larvae)

Ascarids:

Parascaris equorum (luminal L₅ larvae and adults)

Pinworms:

Oxyuris equi (L₄ stage larvae and adults)

Neck threadworms:

Onchocerca spp (microfilariae)

Stomach bots:

Gasterophilus spp (oral and gastric stages)

7. METHOD AND ROUTE(S) OF ADMINISTRATION

Dosage:

One syringe division of paste per 100 kg body weight (based on a recommended dosage of 200 microgram ivermectin per kg body weight). The syringe containing 7.49 g of paste delivers sufficient paste to treat 700 kg of bodyweight at the recommended dose rate.

Administration:

The paste is given by oral route.

To ensure administration of a correct dose, body weight should be determined as accurately as possible. The animal's mouth should be free from food to ensure swallowing. Turn the screw gauge on the syringe plunger to the body weight of the horse. The tip of the syringe barrel should be inserted at the interdental space (the gap between the front and back teeth) and the paste deposited on the base of the tongue. Advance the plunger as far as it will go, depositing the medication on the base of the tongue. Immediately elevate the horse's head for a few seconds to ensure swallowing.

8. WITHDRAWAL PERIOD

Meat and offal: 34 days

Do not use in mares producing milk for human consumption.

9. SPECIAL WARNING(S), IF NECESSARY

Contraindications

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

Do not use in dogs or cats as severe adverse reactions may occur.

Adverse reactions

The effects of GABA agonists are increased by ivermectin

If you notice any serious effects or other effects not mentioned in this leaflet, please inform your veterinary surgeon

Special precautions for use in animals

Some horses with heavy infections of *Onchocerca* spp. microfilariae have experienced oedema and pruritus following treatment. Such reactions are assumed

to be the result of death of large numbers of microfilariae. These signs resolve within a few days but symptomatic treatment may be advisable

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 or misadministration of the product.

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.

Veterinary advice should be given on appropriate dosing programs and stock management to achieve adequate parasite control and reduce the likelihood of anthelmintic resistance developing. In the event that a product is suspected of being ineffective, the animal owner is advised to seek veterinary advice.

Resistance to ivermectin has been reported in *Parascaris equorum*. Therefore, the use of this product should be based on local farm epidemiological information about susceptibility of nematodes and recommendations on how to limit further selection for resistance to anthelmintics

Since ivermectin is highly bound to plasma proteins, special care should be taken in cases of sick animals or in nutritional conditions associated with low plasma protein levels.

Dogs and cats should not be allowed to ingest spilled gel or have access to used packaging due to the potential for adverse effects related to ivermectin toxicity

The product has been formulated for use in horses only. Cats, dogs, especially Collies, Old English Sheepdogs and related breeds or crosses, and also turtles and tortoises may be adversely affected by the concentration of ivermectin in this product if they are allowed to ingest spilled paste or have access to used syringes.

The product can be administered to mares at any stages of pregnancy or lactation.

Mild transitory signs (slowed pupillary light response and depression) have been seen at a higher dose of 1.8 mg/kg (9 times the recommended dose level). Other signs seen at higher doses include mydriasis (dilation of the pupil), ataxia (inco-ordination), tremors (shaking), stupor (dull), coma and death. The less severe signs have been transitory.

Although no antidote has been identified, symptomatic therapy may be beneficial.

User warnings

Do not smoke, eat or drink while handling the product.

Wash hands after use.

This product may cause skin and eye irritation. Therefore, the user should avoid contact of the product with the skin and the eyes. In case of contact, rinse immediately with plenty of water.

In the case of accidental ingestion or eye irritation after contact seek medical advice immediately and show the package insert or the label to the physician.

10. EXPIRY DATE

MM/YYYY

The product is for single use. After use the syringe should be discarded.

11. SPECIAL STORAGE CONDITIONS

Do not store above 25 °C

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

EXTREMELY DANGEROUS TO FISH AND AQUATIC LIFE. Do not contaminate surface waters or ditches with product or used container. Any unused veterinary medicinal product or waste materials derived from such veterinary medicinal products should be disposed of in accordance with local requirements.

13. THE WORDS “FOR ANIMAL TREATMENT ONLY” AND CONDITIONS OR RESTRICTIONS REGARDING SUPPLY AND USE, if applicable

For animal treatment only

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

Keep out of the reach and sight of children

15. NAME AND ADDRESS OF THE MARKETING AUTHORISATION HOLDER

Marketing authorisation holder

ACME DRUGS s.r.l.
Via Portella della Ginestra 9
42025 Cavriago
Italy

Manufacturer for the batch release:

ACME Drugs S.r.l.
Via Portella della Ginestra, 9, Zona Industriale Corte Tegge
42025 CAVRIAGO (RE)
Italy

16. MARKETING AUTHORISATION NUMBER

Veterinary Medicinal Product authorised for use in UK and IE.

UK ONLY

POM-
VPS

To be supplied only on a veterinary prescription'

Vm 61301/5000

Dispose of any unused product and empty containers in accordance with guidance from your local waste regulation authority

17. MANUFACTURER'S BATCH NUMBER

BN XXXX

18. OTHER INFORMATION

Date of approval of the text: August 2025

Gavin Hall
Approved: 29 August 2025