PARTICULARS TO APPEAR ON THE OUTER PACKAGE CARTON

1. NAME OF THE VETERINARY MEDICINAL PRODUCT

Vitofyllin 50 mg film-coated tablets for dogs

2. STATEMENT OF ACTIVE AND OTHER SUBSTANCES

Active substance:
Each tablet contains 50 mg of Propentofylline

3. PACKAGE SIZE

56 tablets
140 tablets

4. TARGET SPECIES

Dogs

5. INDICATION(S)

(*Indication generally to be included only for medicinal products not subject to medicinal prescription*).

6. ROUTE(S) OF ADMINISTRATION

For oral administration.

7. WITHDRAWAL PERIOD

8. EXPIRY DATE

EXP {month/year}
Shelf life of divided tablet portions: 72 hours

9. SPECIAL STORAGE PRECAUTIONS

Divided tablets should be stored in the blister packs.
Keep the blister packs in the outer carton. Store in a dry place.
Store in the original blister package.

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 REACH AND SIGHT OF CHILDREN”

Keep out of the sight and reach of children.

13. NAME AND ADDRESS OF THE MARKETING AUTHORISATION HOLDER

WDT- Wirtschaftsgenossenschaft deutscher Tierärzte eG
Siemensstr. 14
30827 Garbsen
Lower Saxony
Germany

14. MARKETING AUTHORISATION NUMBER

Vm 32829/5000

15. BATCH NUMBER

Lot {number}

16. SPECIAL WARNING(S), IF NECESSARY

Not applicable.

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

Disposal: Read package leaflet

18. CONDITIONS OR RESTRICTIONS REGARDING SUPPLY AND USE IF APPLICABLE

POM-V (To be supplied only on veterinary prescription).

MINIMUM PARTICULARS TO APPEAR ON BLISTERS OR STRIPS
BLISTERS

1. NAME OF THE VETERINARY MEDICINAL PRODUCT

Vitofyllin 50 mg film-coated tablets for dogs

2. QUALITATIVE AND QUANTITATIVE PARTICULARS OF THE ACTIVE SUBSTANCE(S)

Propentofylline	50.00 mg/tablet

3. BATCH NUMBER

Lot {number}

4. EXPIRY DATE

<EXP {month/year}>

5. NAME OF THE MARKETING AUTHORISATION HOLDER

WDT- Wirtschaftsgenossenschaft deutscher Tierärzte eG

6. THE WORDS “FOR ANIMAL TREATMENT ONLY”

For animal treatment only.

PACKAGE LEAFLET

1. NAME OF THE VETERINARY MEDICINAL PRODUCT

Vitofyllin 50 mg film-coated tablets for dogs

2. COMPOSITION

Active substance:
Each tablet contains 50 mg of propentofylline.

Excipients:

Ferric Oxide, yellow, (E 172)	0.075 mg/tablet
Titanium Dioxide, (E171)	0.215 mg/tablet

Film-coated tablets.
Yellow, round, convex tablets with cross breakline on one side and imprinting “50” on the other side.
The tablet can be divided into 2 or 4 equal parts.

3. TARGET SPECIES

Dogs.

4. INDICATION FOR USE

For the improvement of peripheral and cerebral vascular blood circulation. For improvement in dullness, lethargy and overall demeanour in dogs.

5. CONTRAINDICATIONS

Refer to section 6. Special warnings, subsection Pregnancy and lactation.
Do not use in pregnant or lactating bitches or breeding animals.
Do not use in dogs weighing less than 2.5 kg.
Do not use in cases of hypersensitivity to the active substance or to any of the excipients.

6. SPECIAL WARNING(S)

Special precautions for safe use in the target species:
Specific diseases (e.g. kidney disease) should be treated accordingly.
Consideration should be given to rationalising the medication of dogs already receiving treatment for congestive heart failure or bronchial disease.
In the case of renal failure, the dose should be reduced.

Special precautions to be taken by the person administering the veterinary medicinal product to animals:
Care should be taken to avoid accidental ingestion.
In case of accidental ingestion, seek medical advice immediately and show the package leaflet or the label to the physician.
Wash hands after use.

Pregnancy and lactation:
The safety of the product has not been established during pregnancy and lactation. Do not use in pregnant or lactating bitches or breeding animals.

Overdose:
Excitation, tachycardia, hypotension, reddening of mucous membranes and vomiting. The withdrawal of the treatment leads to a spontaneous remission of these signs.

7. ADVERSE EVENTS

Dogs:

Rare (1 to 10 animals / 10,000 animals treated)	Allergic skin reactions*, vomiting*, cardiac disorder*

*In these cases, the treatment should be stopped.

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.
UK national contact details: national system details. https://www.gov.uk/report-veterinary-medicine-problem

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

The basic dosage is 6-10 mg propentofylline/kg bodyweight daily, divided into two 3-5 mg/kg doses as follows:

Body weight (kg)	**Tablets**		**Daily total tablets**	**Daily total dose (mg/kg)**
	am	**pm**		
2.5 - 4 kg	¼	¼	½	6.3 - 10.0
5 - 7 kg	½	½	1	7.1 - 10.0
8 – 9 kg	¾	¾	1 ½	8.3 - 9.4
10 - 15 kg	1	1	2	6.7 – 10.0
16 - 25 kg	1½	1½	3	6.0 – 9.4
26 - 33 kg	2	2	4	6.1- 7.7

To ensure administration of the correct dose, the body weight of the animal should be determined before treatment.
Dogs of more than 20 kg can be given Vitofyllin 100 mg film-coated tablets for dogs.

9. ADVICE ON CORRECT ADMINISTRATION

The tablets can be administered directly onto the back of the dog's tongue or can be mixed in a small ball of food and should be administered at least 30 minutes before feeding.

10. WITHDRAWAL PERIOD(S)

Not applicable.

11. SPECIAL STORAGE PRECAUTIONS

Keep out of the sight and reach children.
Store in the original blister package.
Keep the blister packs in the outer carton.
Store in a dry place.
Unused divided tablets should be returned to the blister pack.
Shelf life of divided tablet portions: 72 hours.
Do not use this veterinary medicinal product after the expiry date which is stated on the carton and blister after Exp.
The expiry date refers to the last day of that month.

12. SPECIAL PRECAUTIONS FOR DISPOSAL

Medicines should not be disposed of via wastewater.
Any unused veterinary medicinal product or waste materials derived from such veterinary medicinal product should be disposed of in accordance with local requirements.

13. CLASSIFICATION OF VETERINARY MEDICINAL PRODUCTS

Veterinary medicinal product subject to prescription.

14. MARKETING AUTHORISATION NUMBERS AND PACK SIZES

Vm 32829/5000

Polyvinylchloride– PolyVinylidene dichloride /Aluminium blister with 14 tablets, in a cardboard box containing 4 blisters (56 tablets).

Polyvinylchloride– PolyVinylidene dichloride /Aluminium blister with 14 tablets, in a cardboard box containing 10 blisters (140 tablets).

Not all pack sizes may be marketed.

15. DATE ON WHICH THE PACKAGE LEAFLET WAS LAST REVISED

January 2023
Find more product information by searching for the ‘Product Information Database’ or ‘PID’ on www.gov.uk

16. CONTACT DETAILS

Marketing authorisation holder:
WDT- Wirtschaftsgenossenschaft deutscher Tierärzte eG
Siemensstr. 14
30827 Garbsen
Lower Saxony
Germany

Manufacturer responsible for batch release:
Artesan Pharma GmbH & Co. KG
Wendlandstraße 1, D – 29439 Lüchow
Germany

17. OTHER INFORMATION

Propentofylline has been shown to increase blood flow, particularly of the heart and skeletal muscle. It also increases the blood flow of the brain and therefore its oxygen supply, without increasing the brain's glucose demand. It has a modest positive chronotropic effect and a marked positive ionotropic effect. In addition, it has been shown to have an anti-arrhythmic effect in dogs with myocardial ischemia and a bronchodilator action equivalent to that of aminofylline.
Propentofylline inhibits platelet aggregation and improves the flow properties of erythrocytes.
It has a direct effect on the heart and reduces peripheral vascular resistance thereby lowering cardiac load.

Propentofylline may increase willingness to exercise and exercise tolerance, particularly in older dogs.

Approved 12 January 2023