

**PARTICULARS TO APPEAR ON THE OUTER PACKAGE**

**NATURE/TYPE:** Cardboard box

**1. NAME OF THE VETERINARY MEDICINAL PRODUCT**

Corvental-D 500 mg Hard Capsules

**2. STATEMENT OF ACTIVE SUBSTANCES**

**Active ingredient:** Each capsule contains 500 mg Theophylline in a sustained release form.

**3. PHARMACEUTICAL FORM**

Hard capsules

**4. PACKAGE SIZE**

100 capsules  
500 mg

**5. TARGET SPECIES**

Dog

**6. INDICATION(S)**

For the treatment of bronchitis and congestive heart failure in the dog.

**7. METHOD AND ROUTE(S) OF ADMINISTRATION**

**Dose:** 20 mg per kg bodyweight to be administered orally once daily.

**8. WITHDRAWAL PERIOD(S)**

**9. SPECIAL WARNING(S), IF NECESSARY**

See package leaflet.

**10. EXPIRY DATE**

USE BY END: {month/year}

**11. SPECIAL STORAGE CONDITIONS**

Do not store above 30°C.  
Store in a dry place.  
Keep blisters in outer carton.

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

See package leaflet.

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

For animal treatment only. To be supplied only on veterinary prescription.

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

Keep out of reach and sight of children.

**15. NAME AND ADDRESS OF THE MARKETING AUTHORISATION HOLDER**

Elanco GmbH  
Heinz-Lohmann Strasse 4  
Groden  
D-27472 Cuxhaven  
Germany

**16. MARKETING AUTHORISATION NUMBER(S)**

Vm 52127/3012

POM-V

**17. MANUFACTURER’S BATCH NUMBER**

BN: {number}

**MINIMUM PARTICULARS TO APPEAR ON BLISTERS OR STRIPS**

**NATURE/TYPE: Blister**

**1. NAME OF THE VETERINARY MEDICINAL PRODUCT**

Corvental-D Hard Capsules Theophylline 500 mg

**2. NAME OF THE MARKETING AUTHORISATION HOLDER**

Elanco GmbH

**3. EXPIRY DATE**

{month/year}

**4. BATCH NUMBER**

{number}

**5. THE WORDS “FOR ANIMAL TREATMENT ONLY”**

FOR ANIMAL TREATMENT ONLY

## **B. PACKAGE LEAFLET**

**PACKAGE LEAFLET:  
Corvental-D Hard Capsules**

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

Marketing authorisation holder:

Elanco GmbH  
Heinz-Lohmann Strasse 4  
Groden  
D-27472 Cuxhaven  
Germany

Manufacturer responsible for batch release:

Swiss Caps GmbH  
Grassingerstraße 9  
83043 Bad Aibling  
Germany

**2. NAME OF THE VETERINARY MEDICINAL PRODUCT**

Corvental-D Capsules 100 mg Hard Capsules  
Corvental-D Capsules 200 mg Hard Capsules  
Corvental-D Capsules 500 mg Hard Capsules

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

Corvental-D Hard Capsules are hard gelatin capsules containing 100 mg, 200 mg or 500 mg Theophylline in a sustained release “Divido” presentation. The capsules are coloured and sized, according to active ingredient content.

Corvental-D Capsules 100 mg Hard Capsules: blue/white, size 3, “Th100” written in black.

Corvental-D Capsules 200 mg Hard Capsules: opaque green/transparent green, size 2, “Th200” written in white.

Corvental-D Capsules 500 mg Hard Capsules: opaque green/transparent green, size 0/elongated, “Th500” written in white.

**4. INDICATION(S)**

For the treatment of bronchitis and congestive heart failure in the dog.

**5. CONTRAINDICATIONS**

Do not use in dogs with a known history of epileptiform seizures as convulsions have been reported in patients on theophylline treatment, often with no preceding signs of toxicity and in otherwise apparently normal animals.

Concurrent use of beta sympathomimetics is contra-indicated, as additive or synergistic interactions resulting in exaggerated side effects may result.

## **6. ADVERSE REACTIONS**

The following side effects have been reported; restlessness, agitation, excitement, vomiting, diarrhoea, polydipsia, sedation, reduced appetite and polyuria. If vomiting occurs the dose should be reduced or the treatment discontinued.

If signs of CNS excitement occur (twitching, restlessness or convulsions) discontinue treatment immediately.

## **7. TARGET SPECIES**

Dog

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

20 mg per kg bodyweight to be administered orally once daily.

## **9. ADVICE ON CORRECT ADMINISTRATION**

For animal treatment only.

Care should be taken to ensure that dogs are weighed carefully and accurately, and the dose does not exceed 20 mg/kg bodyweight.

## **10. WITHDRAWAL PERIOD(S)**

## **11. SPECIAL STORAGE PRECAUTIONS**

Keep out of reach and sight of children.

Do not store above 30°C.

Store in a dry place. Keep blisters in outer carton.

Do not use after the expiry date stated on the carton or blister after EXP.

## **12. SPECIAL WARNING(S)**

Special precautions for use in animals:

Theophylline should be used with caution in patients with liver disease.

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

In the case of accidental ingestion, seek medical advice immediately and show the package leaflet or the label to the physician.

Pregnancy and lactation:

When theophylline is prescribed to pregnant bitches, the risk benefit of the treatment should be assessed. Only small amounts of theophylline are excreted in milk.

Interaction with other medicinal products and other forms of interaction:

Plasma theophylline levels may increase in patients under treatment with macrolide & fluoroquinolone antibiotics such as erythromycin & enrofloxacin, and decrease in patients receiving phenobarbitone or phenytoin.

Theophylline may reduce the convulsive threshold in patients receiving ketamine.

Administration of theophylline shortly before halothane anaesthesia may result in arrhythmogenic effects.

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

Dispose of used packaging in the household refuse. Unused product should be returned to the veterinary surgeon.

**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](http://www.gov.uk).

**15. OTHER INFORMATION**

**POM-V**

To be supplied only on veterinary prescription only.

Packs containing 100 capsules are available for all 3 strengths.

The capsules are packed in blister strips.

100 mg capsules - Vm 52127/3010

200 mg capsules - Vm 52127/3011

500 mg capsules - Vm 52127/3012

Approved 28 March 2025

*Gavin Hall*