**Source URL:** https://medicines.health.europa.eu/veterinary/en/600000063423

# Luminal vet, 100 mg tablets for dogs

Authorised

Phenobarbital

# Product identification

#### **Medicine name:**

Luminal vet, 100 mg tablets for dogs Luminal vet 100 mg Tabletten für Hunde

#### **Active substance:**

Phenobarbital

## **Target species:**

Dog

#### Route of administration:

Oral use

# **Product details**

# **Active substance and strength:**

Phenobarbital 100.00 milligram(s) / 1.00 Tablet

#### **Pharmaceutical form:**

**Tablet** 

#### **Anatomical therapeutic chemical veterinary (ATCvet) codes:**

ON03AA02

#### Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

#### **Authorisation status:**

Valid

#### **Authorised in:**

Germany

## Package description:

(ID7) 120 Tablet: unspecified outer container with 1 Container (High Density

PolyEthylene) with 120 Tablet

(ID6) 100 Tablet: unspecified outer container with 1 Container (High Density

PolyEthylene) with 100 Tablet

(ID5) 60 Tablet: unspecified outer container with 1 Container (High Density

PolyEthylene) with 60 Tablet

(ID4) 50 Tablet: unspecified outer container with 1 Container (High Density

PolyEthylene) with 50 Tablet

(ID3) 180 Tablet: unspecified outer container with 6 Bottle (Glass) each with 30 Tablet

(ID2) 90 Tablet: unspecified outer container with 3 Bottle (Glass) each with 30 Tablet

(ID1) 30 Tablet: unspecified outer container with 1 Bottle (Glass) with 30 Tablet

# Additional information

# **Entitlement type:**

Marketing Authorisation

# Legal basis of product authorisation:

Well-established use application (Article 13a of Directive No 2001/82/EC)

# Marketing authorisation holder:

Desitin Arzneimittel GmbH

# Marketing authorisation date:

2/11/2010

## Manufacturing sites for batch release:

Desitin Arzneimittel GmbH

## **Responsible authority:**

Federal Office Of Consumer Protection And Food Safety

#### **Authorisation number:**

401354.00.00

## Date of authorisation status change:

8/08/2016

#### **Reference member state:**

Germany

#### **Procedure number:**

DE/V/0141/002

#### **Concerned member states:**

Austria Belgium Denmark Netherlands Portugal Spain Sweden

To consult adverse reactions on veterinary medicinal products please go to <a href="https://www.adrreports.eu/vet">www.adrreports.eu/vet</a>

# **Documents**

Summary of Product Characteristics

English (RTF)

Published on: 18/12/2024

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