

# Euthasol vet. 400 mg/ml, solution for injection

Authorised

- Pentobarbital sodium

## Product identification

**Medicine name:**

Euthasol vet. 400 mg/ml, solution for injection

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**Active substance:**

Pentobarbital sodium

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**Target species:**

Cattle

Dog

Goat

Sheep

Horse (non food-producing)

Cat

Mink

Chinchilla

Gerbil

Guinea pig

Hamster

Mouse

Rat

Rabbit (non food-producing)

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**Route of administration:**

Intracardiac use  
Intravenous use

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## Product details

**Active substance and strength:**

Pentobarbital sodium  
400.00 milligram(s) / 1.00 millilitre(s)

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**Pharmaceutical form:**

Solution for injection

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

QN51AA01

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

France

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**Available in:**

France

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**Package description:**

100 ml colourless type I glass vial with a light grey bromobutyl rubber stopper and an aluminium cap.

250 ml colourless type I glass vial with a dark grey bromobutyl rubber stopper and an aluminium cap.

100 ml colourless type II glass vial with a light grey bromobutyl rubber stopper and an aluminium cap. [HISTORICAL]

250 ml colourless type II glass vial with a dark grey bromobutyl rubber stopper and an aluminium cap. [HISTORICAL]

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Hybrid application (Article 13(3) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Le Vet. B.V.

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**Marketing authorisation date:**

15/12/2011

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**Manufacturing sites for batch release:**

Produlab Pharma B.V.

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**Responsible authority:**

French Agency For Food, Environmental And Occupational Health & Safety

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**Authorisation number:**

FR/V/9334132 5/2011

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**Date of authorisation status change:**

30/11/2016

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**Reference member state:**

Ireland

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**Procedure number:**

IE/V/0618/001

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**Concerned member states:**

Austria Belgium Denmark Estonia Finland France Greece Iceland Italy  
Latvia Lithuania Luxembourg Norway Poland Portugal Romania Spain  
Sweden United Kingdom (Northern Ireland)

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To consult adverse reactions on veterinary medicinal products please go to [www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Summary of Product Characteristics

English (PDF)

Published on: 16/01/2026

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Combined File of all Documents

Package Leaflet and Labelling

This document does not exist in this language (English). You can find it in another language below.