

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

Authorised

- Pentobarbital sodium

## Product identification

### Medicine name:

Euthasol vet. 400 mg/ml, solution for injection

EUTHASOL 400 mg/ml SOLUCION INYECTABLE

### Active substance:

Pentobarbital sodium

### 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:**

Spain

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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:**

9/02/2012

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**Manufacturing sites for batch release:**

Produlab Pharma B.V.

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**Responsible authority:**

Spanish Agency Of Medicines And Medical Devices

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**Authorisation number:**

2455 ESP

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**Date of authorisation status change:**

7/03/2019

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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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### Package Leaflet

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

### Labelling

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

### Combined File of all Documents