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

Authorised

- Pentobarbital sodium

## Product identification

**Medicine name:**
Euthasol vet. 400 mg/ml, solution for injection

**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)

**Route of administration:**
Intracardiac use
Intravenous use

## Product details

**Active substance and strength:**
Pentobarbital sodium
400.00 milligram(s) / 1.00 millilitre(s)

**Pharmaceutical form:**
Solution for injection

**Anatomical therapeutic chemical veterinary (ATCvet) codes:**
QN51AA01

**Legal status of supply:**
Veterinary medicinal product subject to veterinary prescription

**Authorisation status:**
Valid

**Authorised in:**
Norway

**Available in:**
Norway

**Package description:**
250 ml colourless type I glass vial with a dark grey bromobutyl rubber stopper and an aluminium cap.
100 ml colourless type I glass vial with a light 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]

## Additional information

**Entitlement type:**
Marketing Authorisation

**Legal basis of product authorisation:**
Hybrid application (Article 13(3) of Directive No 2001/82/EC)

**Marketing authorisation holder:**
Le Vet. B.V.

**Marketing authorisation date:**
10/02/2012

**Manufacturing sites for batch release:**
Produlab Pharma B.V.

**Responsible authority:**
Norwegian Medical Products Agency

**Authorisation number:**
10-7771

**Date of authorisation status change:**
10/02/2012

**Reference member state:**
Ireland

**Procedure number:**
IE/V/0618/001

**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)

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

## Documents

Summary of Product Characteristics

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

Combined File of all Documents

Package Leaflet

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