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# Euthasol 500 mg/ml solution for injection

Authorised

- Pentobarbital sodium

## Product identification

**Medicine name:**
Euthasol 500 mg/ml solution for injection

**Active substance:**
Pentobarbital sodium

**Target species:**
Cattle
Dog
Goat (adult female)
Sheep
Horse
Cat
Rabbit
Mink
Rodents
Pig

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

## Product details

**Active substance and strength:**
Pentobarbital sodium
500.00 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:**
Belgium

**Package description:**
12 Clear Type I glass vials containing 100 ml, closed with a bromobutyl rubber stopper and aluminium cap in a carton box.
12 Polypropylene vials containing 250 ml, closed with a bromobutyl rubber stopper and aluminium cap in a carton box.
12 Clear Type I glass vials containing 250 ml, closed with a bromobutyl rubber stopper and aluminium cap in a carton box.
1 Polypropylene vial containing 100 ml, closed with a bromobutyl rubber stopper and aluminium cap in a carton box.
1 Clear Type I glass vial containing 250 ml, closed with a bromobutyl rubber stopper and aluminium cap in a carton box.
1 Clear Type I glass vial containing 100 ml, closed with a bromobutyl rubber stopper and aluminium cap in a carton box.
12 Polypropylene vials containing 100 ml, closed with a bromobutyl rubber stopper and aluminium cap in a carton box.
1 Polypropylene vial containing 250 ml, closed with a bromobutyl rubber stopper and aluminium cap in a carton box.

## 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. Beheer B.V.

**Marketing authorisation date:**
18/10/2017

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

**Responsible authority:**
Federal Agency For Medicines And Health Products

**Authorisation number:**
This information is not available for this product.

**Date of authorisation status change:**
24/06/2020

**Reference member state:**
Netherlands

**Procedure number:**
NL/V/0320/001

**Concerned member states:**
Austria Belgium Bulgaria Croatia Cyprus Czechia Denmark Estonia Finland France Greece Hungary Iceland Ireland Italy Latvia Lithuania Luxembourg Malta Norway Poland Portugal Romania Slovakia Slovenia Spain Sweden United Kingdom (Northern Ireland)

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

## Documents

Combined File of all Documents

Summary of Product Characteristics

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

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.