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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:
Netherlands

Available in:
Netherlands

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:
30/05/2017

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

Responsible authority:
Medicines Evaluation Board

Authorisation number:
REG NL 119241

Date of authorisation status change:
27/01/2022

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)

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Documents

Combined File of all Documents

English (PDF)
Published on: 20/02/2025
Updated on: 14/03/2026
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Summary of Product Characteristics

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Package Leaflet

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