Euthasol 500 mg/ml solution for injection

Authorised

- Pentobarbital sodium

Product identification

Medicine name:
Euthasol 500 mg/ml solution for injection
Repose 500 mg/ml ενέσιμο διάλυμα

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:
Greece

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:
15/08/2017

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

Responsible authority:
National Organization For Medicines

Authorisation number:
68619/22-06-2022/K-0222501

Date of authorisation status change:
15/08/2017

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.