

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:

Bulgaria

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:

8/06/2017

Manufacturing sites for batch release:

Produlab Pharma B.V.

Responsible authority:

Bulgarian Food Safety Authority

Authorisation number:

0022-2741

Date of authorisation status change:

8/06/2017

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

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.

Labelling

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