

Euthasol vet. 400 mg/ml, solution for injection

Authorised

- Pentobarbital sodium

Product identification

Medicine name:

Euthasol vet. 400 mg/ml, solution for injection
EUTHASOL 400 mg/ml SOLUCION INYECTABLE

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:

Spain

Package description:

100 ml colourless type I glass vial with a light grey bromobutyl rubber stopper and an aluminium cap.

250 ml colourless type I glass vial with a dark 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:

9/02/2012

Manufacturing sites for batch release:

Produlab Pharma B.V.

Responsible authority:

Spanish Agency Of Medicines And Medical Devices

Authorisation number:

2455 ESP

Date of authorisation status change:

7/03/2019

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

English (PDF)

Published on: 16/01/2026

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Package Leaflet

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Labelling

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