Euthasol vet. 400 mg/ml, solution for injection

Authorised

- Pentobarbital sodium

Product identification

Medicine name:
Euthasol vet. 400 mg/ml, solution for injection
Euthasol Vet 400 mg/ml Oplossing voor injectie
Euthasol Vet 400 mg/ml Solution injectable
Euthasol Vet 400 mg/ml Injektionslösung

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

Withdrawal period by route of administration:
Intracardiac use:

- **Cattle**
- **Dog**
- **Goat**
- **Sheep**
- **Horse (non food-producing)**
- **Cat**
- **Mink**
- **Chinchilla**
- **Gerbil**
-

Guinea pig

- **Hamster**
- **Mouse**
- **Rat**
- **Rabbit (non food-producing)**

Intravenous use:

- **Cattle**
- **Dog**
- **Goat**
- **Sheep**
- **Horse (non food-producing)**
- **Cat**
- **Mink**
- **Chinchilla**
- **Gerbil**
- **Guinea pig**
- **Hamster**
- **Mouse**
- **Rat**

- **Rabbit (non food-producing)**

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:
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:
Dechra Regulatory B.V.

Marketing authorisation date:
18/10/2012

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

Responsible authority:
Federal Agency For Medicines And Health Products

Authorisation number:
BE-V428872

Date of authorisation status change:
18/10/2012

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: 28/01/2022
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Package Leaflet

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