

## Euthanimal 20%

Authorised

- Pentobarbital sodium

### Product identification

**Medicine name:**

Euthanimal 20%

Euthanimal 20%, 200 mg/ml ενέσιμο διάλυμα

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**Active substance:**

Pentobarbital sodium

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**Target species:**

Cattle

Dog

Goat (adult female)

Sheep

Horse

Cat

Pig

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**Route of administration:**

Intracardiac use

Intravenous use

### Product details

**Active substance and strength:**

Pentobarbital sodium  
200.00 milligram(s) / 1.00 millilitre(s)

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**Pharmaceutical form:**

Solution for injection

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QN51AA01

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Cyprus

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**Package description:**

6 vials of 250 ml, type II glass injection vial with a bromobutylrubber stopper and aluminium cap in polystyrene box.

12 vials of 100 ml, type II glass injection vial with a bromobutylrubber stopper and aluminium cap in polystyrene box.

1 vial of 250 ml, type II glass injection vial with a bromobutylrubber stopper and aluminium cap in carton box.

1 vial of 100 ml, type II glass injection vial with a bromobutylrubber stopper and aluminium cap in carton box.

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Alfasan Nederland B.V.

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**Marketing authorisation date:**

29/07/2014

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**Manufacturing sites for batch release:**

Alfasan Nederland B.V.

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**Responsible authority:**

Veterinary Services, Ministry Of Agriculture, Natural Resources And Environment

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**Authorisation number:**

CY00466V

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**Date of authorisation status change:**

1/01/2018

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**Reference member state:**

Netherlands

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**Procedure number:**

NL/V/0177/001

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**Concerned member states:**

Belgium Bulgaria Cyprus Czechia Denmark Estonia Finland Germany  
Greece Hungary Ireland Italy Latvia Lithuania Malta Poland Portugal  
Romania Slovenia Spain Sweden United Kingdom (Northern Ireland)

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To consult adverse reactions on veterinary medicinal products please go to  
[www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Combined File of all Documents

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