

Euthanimal 40%

Authorised

- Pentobarbital sodium

Product identification

Medicine name:

Euthanimal 40%

Euthanimal 40%, 400 mg / ml solução injetável

Active substance:

Pentobarbital sodium

Target species:

Cattle

Dog

Goat (adult female)

Sheep

Horse

Cat

Pig

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 (adult female)

-

Sheep

-

Horse

-

Cat

-

Pig

Intravenous use:

-

Cattle

-

Dog

-

Goat (adult female)

-

Sheep

-

Horse

-

Cat

-

Pig

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QN51AA01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Portugal

Package description:

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

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.

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

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

Marketing authorisation holder:

Alfasan Nederland B.V.

Marketing authorisation date:

27/09/2013

Manufacturing sites for batch release:

Alfasan Nederland B.V.

Responsible authority:

Directorate General For Food And Veterinary

Authorisation number:

685/02/13RFVPT

Date of authorisation status change:

11/07/2024

Reference member state:

Netherlands

Procedure number:

NL/V/0177/002

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)

To consult adverse reactions on veterinary medicinal products please go to
www.adrreports.eu/vet

Documents

Combined File of all Documents

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