Euthanimal 20%

Authorised

- Pentobarbital sodium

Product identification

Medicine name:
Euthanimal 20%
Euthanimal 20%, 200 mg/ml raztopina za injiciranje za govedo, konje, prašiče, koze, ovce, pse, mačke, kunce, morske prašičke, miši, podgane, hrčke, piščance, golobe, race, majhne okrasne ptice, kače, želve, kuščarje in žabe

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
200.00 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:
Slovenia

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.

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:
21/06/2013

Manufacturing sites for batch release:
Alfasan Nederland B.V.

Responsible authority:
Agency For Medicinal Products And Medical Devices Of The Republic Of Slovenia

Authorisation number:
MR/V/0434/001

Date of authorisation status change:
21/06/2013

Reference member state:
Netherlands

Procedure number:
NL/V/0177/001

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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