Euthanimal 20%, 200 mg/ml Solution for Injection

Authorised

- Pentobarbital sodium

Product identification

Medicine name:
Euthanimal 20%
Euthanimal 20%, 200 mg/ml Solution for Injection

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:
United Kingdom (Northern Ireland)

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:
2/10/2013

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

Responsible authority:
The Veterinary Medicines Directorate

Authorisation number:
Vm 36408/3050

Date of authorisation status change:
10/10/2024

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)

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