PropoFlo Plus, 10 mg/ml, Emulsion for Injection for Dogs and Cats

Authorised

- Propofol

Product identification

Medicine name:
PropoFlo Plus, 10 mg/ml, Emulsion for Injection for Dogs and Cats

Active substance:
Propofol

Target species:
Dog
Cat

Route of administration:
Intravenous use

Product details

Active substance and strength:
Propofol
10.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Emulsion for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QN01AX10

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
United Kingdom (Northern Ireland)

Available in:
United Kingdom (Northern Ireland)

Package description:
box containing 1 vial of 50 ml
box containing 5 vials of 20 ml

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:
Zoetis Belgium S.A.

Marketing authorisation date:
15/12/2010

Manufacturing sites for batch release:
Fresenius Kabi AB

Responsible authority:

The Veterinary Medicines Directorate

Authorisation number:
Vm 60021/3045

Date of authorisation status change:
2/12/2024

Reference member state:
Spain

Procedure number:
ES/V/0324/001

Concerned member states:
Austria Belgium Denmark Finland France Germany Ireland Italy Luxembourg Netherlands Norway Portugal 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

Summary of Product Characteristics

eu-PUAR-propoflo-plus--10mg-ml--emulsion-for-injection-en.pdf