Vetofol 10 mg/ml Emulsion for Injection for Cats and Dogs

Authorised

- Propofol

Product identification

Medicine name:
Vetofol 10 mg/ml Emulsion for Injection for Cats and Dogs
Vetofol 10 mg/ml Emulsion for Injection for Cats and Dogs

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:
Ireland

Package description:
20ml type I clear glass vial sealed with bromobutyl bung and aluminium seal.Available in carton of 1 x 20ml
50ml type I clear glass vial sealed with bromobutyl bung and aluminium seal.Available in carton of 1 x 50ml
20ml type I clear glass vial sealed with bromobutyl bung and aluminium seal.Available in carton of 5 x 20ml
50ml type I clear glass vial sealed with bromobutyl bung and aluminium seal.Available in carton of 5 x 50ml

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:
Norbrook Laboratories (Ireland) Limited

Marketing authorisation date:
31/08/2012

Manufacturing sites for batch release:

Norbrook Laboratories Limited
Norbrook Manufacturing Limited

Responsible authority:
Health Products Regulatory Authority

Authorisation number:
VPA22664/109/001

Date of authorisation status change:
31/08/2012

Reference member state:
Ireland

Procedure number:
IE/V/0533/001

Concerned member states:
Denmark France Italy Norway Sweden United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics