

Vetofol 10mg/ml Emulsion for Injection for cats and dogs

Authorised

- Propofol

Product identification

Medicine name:

Vetofol 10mg/ml Emulsion for Injection for cats and dogs

Vetofol 10mg/ml Ενέσιμο γαλάκτωμα για γάτες και σκύλους

Active substance:

Propofol

Target species:

Dog

Cat

Route of administration:

Intravenous use

Product details

Active substance and strength:

Propofol

1.00 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:

Cyprus

Package description:

(ID2) 50 millilitre(s): unspecified outer container with 1 Vial (glass) with 50 millilitre(s), closed with Stopfen (bromobutyl rubber)
(ID1) 20 millilitre(s): unspecified outer container with 1 Vial (glass) with 20 millilitre(s), closed with Stopfen (bromobutyl rubber)

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:

10/04/2011

Manufacturing sites for batch release:

Norbrook Laboratories (Ireland) Limited

Norbrook Laboratories Limited

Responsible authority:

Veterinary Services, Ministry Of Agriculture, Natural Resources And Environment

Authorisation number:

CY00227V

Date of authorisation status change:

4/03/2015

Reference member state:

Germany

Procedure number:

DE/V/0317/001

Concerned member states:

Austria Bulgaria Cyprus Czechia Estonia Finland Latvia Lithuania Portugal
Romania Slovakia Slovenia Spain United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics

English (RTF)

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