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

Authorised

- Propofol

Product identification

Medicine name:
Norofol 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:
United Kingdom (Northern Ireland)

Package description:
(ID1) 20 millilitre(s): Box (cardboard) with 1 Vial (clear glass (Glass typ I)) with 20 millilitre(s), closed with Stopper (bromobutyl rubber) and Cap`` (aluminium)
(ID2) 50 millilitre(s): Box (cardboard) with 1 Vial (clear glass (Glass typ I)) with 50 millilitre(s), closed with Stopfen (bromobutyl rubber) and Cap`` (aluminium)

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 Limited

Marketing authorisation date:
9/01/2009

Manufacturing sites for batch release:
Norbrook Laboratories (Ireland) Limited
Norbrook Laboratories Limited

Responsible authority:
The Veterinary Medicines Directorate

Authorisation number:
Vm 02000/4275

Date of authorisation status change:
2/01/2024

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

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