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PropoFlo Plus, 10mg/ml, Emulsion for Injection

Authorised

- Propofol

Product identification

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

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:
Norway

Available in:
Norway

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 Animal Health ApS

Marketing authorisation date:
27/06/2012

Manufacturing sites for batch release:
Fresenius Kabi AB

Responsible authority:
Norwegian Medical Products Agency

Authorisation number:

11-8713

Date of authorisation status change:
27/06/2012

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

Summary of Product Characteristics

English (PDF)
Published on: 14/03/2026
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Package Leaflet

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Combined File of all Documents

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