

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:

Austria

Available in:

Austria

Package description:

box containing 5 vials of 20 ml

box containing 1 vial of 50 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 Österreich GmbH

Marketing authorisation date:

4/05/2012

Manufacturing sites for batch release:

Fresenius Kabi AB

Responsible authority:

Austrian Agency For Health And Food Safety

Authorisation number:

8-01074

Date of authorisation status change:

4/05/2012

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

Package Leaflet

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Summary of Product Characteristics

English (PDF)

Published on: 6/04/2023

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Labelling

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Combined File of all Documents

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