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:
Belgium

Available in:
Belgium

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 Belgium

Marketing authorisation date:
14/12/2021

Manufacturing sites for batch release:
Fresenius Kabi AB

Responsible authority:
Federal Agency For Medicines And Health Products

Authorisation number:

BE-V420472

Date of authorisation status change:
14/12/2021

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: 6/04/2023
Download

Package Leaflet

This document does not exist in this language (English). You can find it in another language below.

Labelling

This document does not exist in this language (English). You can find it in another language below.

Combined File of all Documents

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