Propodine 10 mg/ml Emulsion for Injection/Infusion for Dogs and Cats

Authorised

- Propofol

Product identification

Medicine name:
Propodine 10 mg/ml Emulsion for Injection/Infusion for Dogs and Cats

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/infusion

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QN01AX10

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Finland

Package description:
Colourless type I glass vials of 100 ml closed with a coated bromobutyl rubber stopper and aluminium cap in a carton box.
Colourless type I glass vials of 20 ml closed with a coated bromobutyl rubber stopper and aluminium cap in a carton box.
Colourless type I glass vials of 50 ml closed with a coated bromobutyl rubber stopper and aluminium cap in a carton box.

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Well-established use application (Article 13a of Directive No 2001/82/EC)

Marketing authorisation holder:
Dechra Regulatory B.V.

Marketing authorisation date:
20/08/2020

Manufacturing sites for batch release:
Corden Pharma S.p.A.

Responsible authority:
Finnish Medicines Agency

Authorisation number:
36146

Date of authorisation status change:
20/08/2020

Reference member state:
Netherlands

Procedure number:
NL/V/0319/001

Concerned member states:
Austria Belgium Bulgaria Croatia Cyprus Czechia Denmark Estonia Finland France Germany Greece Hungary Iceland Ireland Italy Latvia Lithuania Luxembourg Norway Poland Portugal Romania Slovakia Slovenia Spain Sweden United Kingdom (Northern Ireland)

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