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

- Propofol

Authorised

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:

- Poland

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:

- 2/07/2020

Manufacturing sites for batch release:

- Corden Pharma S.p.A.

Responsible authority:

- Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

Authorisation number:

- 2997

Date of authorisation status change:

- 2/07/2020

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

Documents

Product information

Package Leaflet

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Labelling

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