

Fentadon 50 microgram/ml, solution for injection for dogs

Authorised

- Fentanyl citrate

Product identification

Medicine name:

Fentadon 50 microgram/ml, solution for injection for dogs

Active substance:

Fentanyl citrate

Target species:

Dog

Route of administration:

Intravenous use

Product details

Active substance and strength:

Fentanyl citrate

78.50 microgram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection/infusion

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QN02AB03

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Portugal

Available in:

Portugal

Package description:

Vial of uncoloured glass type I, filled with 10 ml. Teflon-coated chlorobutyl rubber stoppers type I secured with aluminium caps.

Vial of uncoloured glass type I, filled with 100 ml. Teflon-coated chlorobutyl rubber stoppers type I secured with aluminium caps.

Vial of uncoloured glass type I, filled with 50 ml. Teflon-coated chlorobutyl rubber stoppers type I secured with aluminium caps.

Vial of uncoloured glass type I, filled with 5 ml. Teflon-coated chlorobutyl rubber stoppers type I secured with aluminium caps.

Vial of uncoloured glass type I, filled with 30 ml. Teflon-coated chlorobutyl rubber stoppers type I secured with aluminium caps.

Vial of uncoloured glass type I, filled with 25 ml. Teflon-coated chlorobutyl rubber stoppers type I secured with aluminium caps.

Vial of uncoloured glass type I, filled with 20 ml. Teflon-coated chlorobutyl rubber stoppers type I secured with aluminium caps.

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:

Eurovet Animal Health B.V.

Marketing authorisation date:

17/02/2012

Manufacturing sites for batch release:

Eurovet Animal Health B.V.

Responsible authority:

Directorate General For Food And Veterinary

Authorisation number:

433/01/12DFVPT

Date of authorisation status change:

24/06/2025

Reference member state:

Netherlands

Procedure number:

NL/V/0155/001

Concerned member states:

Austria Belgium Denmark France Germany Italy Luxembourg Norway
Poland Portugal Spain Sweden United Kingdom (Northern Ireland)

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

Documents

Combined File of all Documents

English (PDF)

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