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
Fentadon vet. 50 mikrogram/ml Injektionsvätska, lösning

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

Withdrawal period by route of administration:

Intravenous use:

- **Dog**

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QN02AB03

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Sweden

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:
21/02/2013

Manufacturing sites for batch release:
Eurovet Animal Health B.V.

Responsible authority:
Medical Products Agency

Authorisation number:
44985

Date of authorisation status change:
21/02/2013

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

Package Leaflet

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

Summary of Product Characteristics

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

Combined File of all Documents

108332 PAR.pdf

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