Insistor 10 mg/ml solution for injection for dogs and cats

Authorised

- Methadone hydrochloride

Product identification

Medicine name:
Insistor 10 mg/ml solution for injection for dogs and cats
Insistor 10 mg/ml Roztwór do wstrzykiwań

Active substance:
Methadone hydrochloride

Target species:
Dog
Cat

Route of administration:
Intramuscular use
Intravenous use
Subcutaneous use

Product details

Active substance and strength:
Methadone hydrochloride
10.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Dog**
- **Cat**

Intravenous use:

- **Dog**
- **Cat**

Subcutaneous use:

- **Dog**
- **Cat**

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QN02AC90

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Poland

Available in:

Poland

Package description:

Clear glass vial with grey, coated chlorobutyl rubber stopper and aluminium pull off cap or aluminium/plastic flip off cap. Package size 1x 5 ml

Clear glass vial with grey, coated chlorobutyl rubber stopper and aluminium pull off cap or aluminium/plastic flip off cap. Package size 5x 10 ml
Clear glass vial with grey, coated chlorobutyl rubber stopper and aluminium pull off cap or aluminium/plastic flip off cap. Package size 5 x 5 ml
Clear glass vial with grey, coated chlorobutyl rubber stopper and aluminium pull off cap or aluminium/plastic flip off cap. Package size 1x 10 ml

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:
Vetviva Richter GmbH

Marketing authorisation date:
25/06/2021

Manufacturing sites for batch release:
Vetviva Richter GmbH

Responsible authority:
Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

Authorisation number:
3106

Date of authorisation status change:
25/06/2021

Reference member state:
Netherlands

Procedure number:
NL/V/0235/001

Concerned member states:
Austria Belgium Bulgaria Croatia Cyprus Czechia Denmark Estonia Finland France Germany Greece Hungary Iceland Ireland Italy Latvia Lithuania 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

Labelling

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Summary of Product Characteristics

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

Package Leaflet

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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