

Narcostart (Sedastart) 1 mg/ml solution for injection for cats and dogs

Authorised

- Medetomidine hydrochloride

Product identification

Medicine name:

Narcostart (Sedastart) 1 mg/ml solution for injection for cats and dogs

Narcostart 1 mg/ml injekčný roztok pre psov a mačky

Active substance:

Medetomidine hydrochloride

Target species:

Dog

Cat

Route of administration:

Intramuscular use

Intravenous use

Product details

Active substance and strength:

Medetomidine hydrochloride

1.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QN05CM91

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Slovakia

Package description:

1 x 1 glass vial with 10 ml. Glass (type I) vials closed with bromobutyl rubber closures secured with aluminium crimp caps.

5 x 1 glass vial with 10 ml. Glass (type I) vials closed with bromobutyl rubber closures secured with aluminium crimp caps.

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:

Le Vet. B.V.

Marketing authorisation date:

7/06/2010

Manufacturing sites for batch release:

Produlab Pharma B.V.

Eurovet Animal Health B.V.

Responsible authority:

Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:

96/018/DC/10-S

Date of authorisation status change:

7/06/2010

Reference member state:

Netherlands

Procedure number:

NL/V/0138/001

Concerned member states:

Austria Belgium Czechia Denmark Finland France Greece Hungary Iceland
Ireland Italy Luxembourg Poland Portugal Slovakia Spain Sweden
United Kingdom (Northern Ireland)

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www.adrreports.eu/vet

Documents

Combined File of all Documents

English (PDF)

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