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 stungulyf, lausn fyrir hunda og ketti.

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:
Iceland

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:
27/12/2012

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

Responsible authority:
Icelandic Medicines Agency

Authorisation number:
IS/2/12/016/01

Date of authorisation status change:
26/10/2015

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)

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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