NARCOSTART

Authorised

- Dexmedetomidine hydrochloride

Product identification

Medicine name:
NARCOSTART

Active substance:
Dexmedetomidine hydrochloride

Target species:
Dog
Cat

Route of administration:
Intravenous use
Intramuscular use
Subcutaneous use
Intramuscular use

Product details

Active substance and strength:
Dexmedetomidine 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:
This information is not available for this product.

Authorisation status:
Valid

Authorised in:
Romania

Package description:
Available only in Romanian
Available only in Romanian

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Complete application (stand-alone) - Directive No 2001/82/EC

Marketing authorisation holder:
Maravet S.R.L.

Marketing authorisation date:
26/02/2015

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

Responsible authority:
Institute For Control Of Biological Products And Veterinary Medicines

Authorisation number:

200099

Date of authorisation status change:
24/08/2025

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

Documents

Combined File of all Documents

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