File downloaded on 2026-10-30
Source URL: https://medicines.health.europa.eu/veterinary/en/600000037119
Narcostart (Sedastart) 1 mg/ml solution for injection for cats and dogs

- Medetomidine hydrochloride

Ima dovoljenje za promet

Informacije o zdravilu

Ime zdravila:
Narcostart (Sedastart) 1 mg/ml solution for injection for cats and dogs
U?inkovina:

- Na voljo samo v English

Ciljne živalske vrste:

- Na voljo samo v Bulgarian Spanish Czech Danish German Estonian Greek English French Italian Latvian Lithuanian Hungarian Dutch Romanian Finnish Swedish Icelandic Norwegian
- Na voljo samo v Bulgarian Spanish Czech Danish German Estonian Greek English French Italian Latvian Lithuanian Hungarian Dutch Romanian Finnish Swedish Icelandic Norwegian

Pot uporabe:

- intramuskularna uporaba
- Intravenska uporaba

Podatki o zdravilu

U?inkovina / Jakost:

- Na voljo samo v English
  1.00
  milligram(s)
  /
  1.00
  millilitre(s)

Farmacevtska oblika:

- Raztopina za injiciranje

Anatomsko-terapevtsko-kemi?na veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):

- QN05CM91

Pravni status za dobavo/izdajo zdravila:

- Zdravilo, ki se izdaja na veterinarski recept

Status dovoljenja za promet z zdravilom:

- Valid

Ima dovoljenje za promet v:

- Na voljo samo v Spanish Czech German Estonian English French Croatian Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian

Na voljo v:

- Spain

Opis ovojnine zdravila:

- Na voljo samo v English
- Na voljo samo v English

Dodatne informacije

Vrsta dovoljenja:

- Na voljo samo v English French Croatian Italian Latvian Finnish Swedish Icelandic Norwegian

Pravna podlaga za izdajo dovoljenja za promet z zdravilom:

- Na voljo samo v English Italian Latvian Norwegian

Imetnik dovoljenja za promet z zdravilom:

- Le Vet. B.V.

Datum dovoljenja za promet z zdravilom:

- 29/09/2010

Proizvodna mesta, odgovorna za sproš?anje serij:

- Produlab Pharma B.V.
- Eurovet Animal Health B.V.

Pristojni organ:

- Spanish Agency Of Medicines And Medical Devices

Številka dovoljenja za promet z zdravilom:

- 2195 ESP

Datum spremembe statusa dovoljenja za promet:

- 7/03/2019

Referen?na država ?lanica:

- Na voljo samo v Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian

Številka postopka:

- NL/V/0138/001

Zadevne države ?lanice:

- Na voljo samo v Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian
- Na voljo samo v Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian
- Na voljo samo v Spanish Czech German Estonian English French Italian Latvian Dutch Portuguese Slovak Swedish Icelandic Norwegian
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- Na voljo samo v Spanish Czech German Estonian English French Croatian Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian
- Na voljo samo v Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian
- Na voljo samo v Estonian English French Lithuanian Portuguese Swedish Icelandic Norwegian

Za poro?ila o domnevnih neželenih u?inkih zdravil za uporabo v veterinaski medicini obiš?ite stran: www.adrreports.eu/vet

Dokumenti

Informacije o zdravilu

Povzetek glavnih zna?ilnosti zdravila

Ta dokument ne obstaja v tem jeziku (slovenš?ina). Lahko ga najdete spodaj v drugem jeziku.
Drugi jeziki (1)
Spanish (PDF)
Objavljeno na dan: 21/06/2024
Prenesi

Navodilo za uporabo

Ta dokument ne obstaja v tem jeziku (slovenš?ina). Lahko ga najdete spodaj v drugem jeziku.
Drugi jeziki (1)
Spanish (PDF)
Objavljeno na dan: 21/06/2024
Prenesi

Ozna?evanje

Ta dokument ne obstaja v tem jeziku (slovenš?ina). Lahko ga najdete spodaj v drugem jeziku.
Drugi jeziki (1)
Spanish (PDF)
Objavljeno na dan: 14/03/2026
Prenesi

Combined File of all Documents

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Drugi jeziki (1)
English (PDF)
Objavljeno na dan: 3/06/2024
Prenesi