File downloaded on 2026-11-14
Source URL: https://medicines.health.europa.eu/veterinary/en/600000035050
Bupredine Multidose 0.3 mg/ml Solution for Injection for Dogs, Cats and Horses

- Buprenorphine hydrochloride

Ima dovoljenje za promet

Informacije o zdravilu

Ime zdravila:
Bupredine Multidose 0.3 mg/ml Solution for Injection for Dogs, Cats and Horses
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
- 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
  0.32
  milligram(s)
  /
  1.00
  millilitre(s)

Farmacevtska oblika:

- Raztopina za injiciranje

Karenca glede na pot uporabe:

- Intravenska uporaba
  - Horse

- All relevant tissues
  no withdrawal period
  Not authorised for use in horses intended for human consumption.

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

- QN02AE01

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 Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian

Opis ovojnine zdravila:

- Na voljo samo v English
- Na voljo samo v English
- Na voljo samo v English
- 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. Beheer B.V.

Datum dovoljenja za promet z zdravilom:

- 11/11/2015

Proizvodna mesta, odgovorna za sproš?anje serij:

- Produlab Pharma B.V.

Pristojni organ:

- Danish Medicines Agency

Številka dovoljenja za promet z zdravilom:

- 54940

Datum spremembe statusa dovoljenja za promet:

- 11/11/2015

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/0314/001

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 Croatian Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian
- Na voljo samo v Spanish Czech German Estonian Greek 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
- Na voljo samo v Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian
- Na voljo samo v Spanish German Estonian English French Italian Dutch Portuguese Slovak Finnish 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 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 Dutch Portuguese Slovak Swedish Icelandic Norwegian
- Na voljo samo v Spanish 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 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 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 Dutch Portuguese Slovak Swedish Icelandic Norwegian
- Na voljo samo v Spanish Czech German Estonian English French Italian Lithuanian Dutch Portuguese Romanian 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 Dutch Portuguese Slovak Swedish Icelandic Norwegian
- 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)
Danish (, INJEKTIONSVÆSKE, OPLØSNING 0,3 MG-ML)
Objavljeno na dan: 11/11/2026
Prenesi

Combined File of all Documents

Ta dokument ne obstaja v tem jeziku (slovenš?ina). Lahko ga najdete spodaj v drugem jeziku.
Drugi jeziki (1)
English (PDF)
Objavljeno na dan: 3/11/2026
Prenesi