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Procamidor Duo 40 mg/ml + 0,036 mg/ml Injektionslösung für Tiere

Ima dovoljenje za promet

- Procaine hydrochloride
- Adrenaline hydrogen tartrate

Informacije o zdravilu

Ime zdravila:
Procamidor Duo 40 mg/ml + 0,036 mg/ml Injektionslösung für Tiere

Učinkovina:
Na voljo samo v English
Na voljo samo v English

Ciljne živalske vrste:
govedo
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:

Perinevralna uporaba

Subkutana uporaba

Podatki o zdravilu

Učinkovina / Jakost:

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

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

Farmacevtska oblika:

Raztopina za injiciranje

Karenca glede na pot uporabe:

Perinevralna uporaba:

- **govedo**
  - Milk. 0 hour
  - Meat and offal. 0 day
- **Sheep**
  - Milk. 0 hour
  - Meat and offal. 0 day
- **Horse**
  - Milk. 0 hour
  - Meat and offal. 0 day
- **Pig**
  - Meat and offal. 0 day

Subkutana uporaba:

- **govedo**
  - Milk. 0 hour
  - Meat and offal. 0 day
- **Sheep**
  - Milk. 0 hour
  - Meat and offal. 0 day
- **Horse**
  - Milk. 0 hour
  - Meat and offal. 0 day
- **Pig**
  - Meat and offal. 0 day

Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):
QN01BA52

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

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Na voljo v:
Germany

Opis ovojnine zdravila:

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:
Vetviva Richter GmbH

Datum dovoljenja za promet z zdravilom:
12/07/2019

Proizvodna mesta, odgovorna za sproščanje serij:
Vetviva Richter GmbH

Pristojni organ:
Federal Office Of Consumer Protection And Food Safety

Številka dovoljenja za promet z zdravilom:
402565.00.00

Datum spremembe statusa dovoljenja za promet:
12/07/2019

Referenčna država članica:
Na voljo samo v Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian

Številka postopka:
AT/V/0018/001

Zadevne države članice:

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
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Na voljo samo v Estonian English French Lithuanian Portuguese Swedish Icelandic Norwegian

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Za poročila o domnevnih neželenih učinkih zdravil za uporabo v veterinaski medicini obiščite stran: www.adrreports.eu/vet

Dokumenti

Combined File of all Documents

Ta dokument ne obstaja v tem jeziku (slovenščina). Lahko ga najdete spodaj v drugem jeziku.

Povzetek glavnih značilnosti zdravila

Ta dokument ne obstaja v tem jeziku (slovenščina). Lahko ga najdete spodaj v drugem jeziku.

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