Butomidor 10 mg/ml Injektionslösung für Pferde, Hunde und Katzen

Pooblaščeno

- Butorphanol tartrate

Product identification

Ime zdravila:
Butomidor 10 mg/ml Injektionslösung für Pferde, Hunde und Katzen
Butomidor, 10 mg/ml, injekcinis tirpalas arkliams, šunims ir katėms

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Učinkovina:
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 Swedish Icelandic Norwegian
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Pot uporabe:
intramuskularna uporaba
Intravenska uporaba
Subkutana uporaba

Product details

Učinkovina / Jakost:

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

Farmacevtska oblika:

Raztopina za injiciranje

Withdrawal period by route of administration:

intramuskularna uporaba:

- **Dog**

Intravenska uporaba:

- **Dog**
- **Horse**
  - Meat and offal. 0 day
  - Milk. 0 hour
- **Cat**

Subkutana uporaba:

- **Dog**
- **Cat**
  - Meat and offal. 0 day
  - Milk. 0 hour

Anatomsko-terapevtsko-kemična veterinarska oznaka (ATCvet):

QN02AF01

Pravni status za dobavo / izdajo zdravila:

Na voljo samo v English French Italian Latvian Portuguese Finnish Swedish Norwegian

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Status dovoljenja:

Valid

Authorised in:

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

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Opis ovojnine:
Na voljo samo v English
Na voljo samo v English
Na voljo samo v English
Na voljo samo v English

Additional information

Entitlement type:
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:
Richter Pharma AG

Marketing authorisation date:
9/12/2010

Proizvodna mesta, odgovorna za sproščanje serij:
Vetviva Richter GmbH

Pristojni organ:
State Food And Veterinary Service

Številka dovoljenja :
LT/2/10/1992/001-004

Datum spremembe statusa dovoljenja:
6/10/2015

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

Številka postopka:

Zadevne države članice:

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

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Na voljo samo v Estonian English French Swedish Icelandic Norwegian

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

Documents

Butomidor 2023-09-07.pdf
