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Catobevit 100 mg/ml + 0,05 mg/ml raztopina za injiciranje za govedo, konje in pse

Ima dovoljenje za promet

- Butafosfan
- Cyanocobalamin

Informacije o zdravilu

Ime zdravila:
Catobevit 100 mg/ml + 0,05 mg/ml raztopina za injiciranje za govedo, konje in pse

Učinkovina:
Na voljo samo v English
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
govedo
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

Subkutana uporaba

Podatki o zdravilu

Učinkovina / Jakost:

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

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

Farmacevtska oblika:

Raztopina za injiciranje

Karenca glede na pot uporabe:

Intravenska uporaba:

- **govedo**
  - Milk. 0 day
  - Meat and offal. 0 day
- **Horse**
  - Milk. 0 day
  - Meat and offal. 0 day

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

QA12CX99

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

Na voljo v:
Slovenia

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:
KRKA tovarna zdravil d.d. Novo mesto

Datum dovoljenja za promet z zdravilom:
12/12/2018

Proizvodna mesta, odgovorna za sproščanje serij:
KRKA tovarna zdravil d.d. Novo mesto
TAD Pharma GmbH

Pristojni organ:
Agency For Medicinal Products And Medical Devices Of The Republic Of Slovenia

Številka dovoljenja za promet z zdravilom:
DC/V/0645/001

Datum spremembe statusa dovoljenja za promet:
12/12/2018

Referenčna država članica:

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

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Številka postopka:

CZ/V/0145/001

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 Spanish Czech German Estonian English French Croatian Italian Dutch Portuguese Slovak 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

Povzetek glavnih značilnosti zdravila

Označevanje

Navodilo za uporabo