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T 61 oplossing voor injectie

Ima dovoljenje za promet

- Embutramide
- Tetracaine hydrochloride
- Mebezonium iodide

Informacije o zdravilu

Ime zdravila:
T 61 oplossing voor injectie

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

Ciljne živalske vrste:
Na voljo samo v Bulgarian Spanish Czech Danish Estonian English Italian Latvian Lithuanian Hungarian Romanian Swedish
Na voljo samo v Bulgarian Spanish Czech Danish German Estonian Greek English Latvian Lithuanian Hungarian Romanian Swedish
okrasna ptica
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 Danish German Estonian Greek English Italian Latvian Lithuanian Hungarian Romanian Swedish Icelandic

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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:
Intrakardialna uporaba
Intravenska uporaba
Intrapulmonalna uporaba

Podatki o zdravilu

Učinkovina / Jakost:
Na voljo samo v English
200.00 milligram(s) / 1.00 millilitre(s)

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

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

Farmacevtska oblika:
Raztopina za injiciranje

Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):
QN51AX50

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:
Netherlands

Opis ovojnine zdravila:
Na voljo samo v Dutch

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 French Italian Latvian Norwegian

Imetnik dovoljenja za promet z zdravilom:
Intervet Nederland B.V.

Datum dovoljenja za promet z zdravilom:
8/09/1992

Proizvodna mesta, odgovorna za sproščanje serij:
Intervet International GmbH

Pristojni organ:
Medicines Evaluation Board

Številka dovoljenja za promet z zdravilom:
REG NL 5356

Datum spremembe statusa dovoljenja za promet:
5/02/2015

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.