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T 61, injekcinis tirpalas

Ima dovoljenje za promet

- Embutramide
- Mebezonium iodide
- Tetracaine hydrochloride

Informacije o zdravilu

Ime zdravila:
T 61, injekcinis tirpalas

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 German Estonian Greek English French Italian Latvian Lithuanian Hungarian Dutch Romanian Finnish Swedish Icelandic Norwegian
Na voljo samo v Bulgarian Spanish Czech Greek English Italian Latvian Lithuanian Hungarian Romanian
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 Estonian Greek English Italian Latvian Lithuanian Hungarian Romanian Swedish Icelandic

Pot uporabe:
Intravenska uporaba

Podatki o zdravilu

Učinkovina / Jakost:

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

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

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

Farmacevtska oblika:

Raztopina za injiciranje

Karenca glede na pot uporabe:

Intravenska uporaba:

- **Dog**
  - All relevant tissues. no withdrawal period

T 61 euthanized animals must be disposed of in accordance with national requirements.

- **Large animals**
  - All relevant tissues. no withdrawal period

Do not use in animals intended for human consumption. T 61 euthanized animals must be disposed of in accordance with national requirements.

Intrapulmonalna uporaba:

- **Dog**
  - All relevant tissues. no withdrawal period

T 61 euthanized animals must be disposed of in accordance with national requirements.

- **Cat**
  - All relevant tissues. no withdrawal period

T 61 euthanized animals must be disposed of in accordance with national requirements.

- **Birds**
  - All relevant tissues. no withdrawal period

Do not use in animals intended for human consumption. T 61 euthanized animals must be disposed of in accordance with national requirements.

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

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Opis ovojnine zdravila:
Na voljo samo v Lithuanian

Dodatne informacije

Vrsta dovoljenja:
Na voljo samo v English French Croatian Italian Latvian Finnish Swedish Icelandic Norwegian

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Pravna podlaga za izdajo dovoljenja za promet z zdravilom:
Na voljo samo v English Italian Latvian Lithuanian Norwegian

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

Datum dovoljenja za promet z zdravilom:
15/04/2002

Proizvodna mesta, odgovorna za sproščanje serij:
Intervet International GmbH

Pristojni organ:
State Food And Veterinary Service

Številka dovoljenja za promet z zdravilom:
Ta podatek za to zdravilo ni na voljo.

Datum spremembe statusa dovoljenja za promet:
15/04/2007

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

Dokumenti

RV1399.pdf