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T 61 solution injectable

Ima dovoljenje za promet

- Embutramide
- Mebezonium iodide
- Tetracaine hydrochloride

Informacije o zdravilu

Ime zdravila:
T 61 solution injectable

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

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

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Pot uporabe:
Intrapulmonalna uporaba
Intrakardialna uporaba
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:
Intrapulmonalna uporaba:

- **Pigeon**
  - Meat and offal. no withdrawal period

Euthanised animals should not be used as food (animal or human). Adequate measures should be taken to ensure that carcasses of animals treated with this product and the by-products of these animals do not enter the food chain and are not used for human or animal consumption.

- **Mink**
  - Meat and offal. no withdrawal period

Euthanised animals should not be used as food (animal or human). Adequate measures should be taken to ensure that carcasses of animals treated with this product and the by-products of these animals do not enter the food chain and are not used for human or animal consumption.

- **okrasna ptica**
  - Meat and offal. no withdrawal period

Euthanised animals should not be used as food (animal or human). Adequate measures should be taken to ensure that carcasses of animals treated with this product and the by-products of these animals do not enter the food chain and are not used for human or animal consumption.

- **Laboratory animals**
  - Meat and offal. no withdrawal period

Euthanised animals should not be used as food (animal or human). Adequate measures should be taken to ensure that carcasses of animals treated with this product and the by-products of these animals do not enter the food chain and are not used for human or animal consumption.

Intravenska uporaba:

- **govedo**
  - Meat and offal. no withdrawal period

Euthanised animals should not be used as food (animal or human). Adequate measures should be taken to ensure that carcasses of animals treated with this product and the by-products of these animals do not enter the food chain and are not used for human or animal consumption.

- **Horse**

- Meat and offal. no withdrawal period

Euthanised animals should not be used as food (animal or human). Adequate measures should be taken to ensure that carcasses of animals treated with this product and the by-products of these animals do not enter the food chain and are not used for human or animal consumption.

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:
Luxembourg

Opis ovojnine zdravila:
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

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

Datum dovoljenja za promet z zdravilom:
5/11/1996

Proizvodna mesta, odgovorna za sproščanje serij:
Intervet International GmbH

Pristojni organ:
Ministry Of Health And Social Security

Številka dovoljenja za promet z zdravilom:
V 416/97/04/0547

Datum spremembe statusa dovoljenja za promet:
5/11/2001

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

Dokumenti

Povzetek glavnih značilnosti zdravila

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