

T 61 Oplossing voor injectie

Atļautas

- Mebezonium iodide
- Embutramide
- Tetracaine hydrochloride

Zāļu identifikācija

Zāļu nosaukums:

T 61 Oplossing voor injectie

Aktīvā viela:

Pieejamas tikai [English](#)

Pieejamas tikai [English](#)

Pieejamas tikai [English](#)

Mērķsugas:

Suns

Kaķis

Ūdele

Dekoratīvie putni

Laboratorijas dzīvnieki

Balodis

Liellops

Zirgs

Lietošanas veids:

Intrapulmonālai lietošanai

intrakardiālai lietošanai

intravenozai lietošanai

Sīkāka informācija par zālēm

Aktīvā viela un stiprums:

Pieejamas tikai [English](#)

50.00 miligrams(i) / 1.00 mililitrs(i)

Pieejamas tikai [English](#)

200.00 miligrams(i) / 1.00 mililitrs(i)

Pieejamas tikai [English](#)

5.00 miligrams(i) / 1.00 mililitrs(i)

Farmaceutiskā forma:

Šķīdums injekcijām

Ierobežojumu periods pēc lietošanas veida:

Intrapulmonālai lietošanai:

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Suns

- Gaļa un blakusprodukti. 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

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Kaķis

- Gaļa un blakusprodukti. no withdrawal period

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Dekoratīvie putni

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Laboratorijas dzīvnieki

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intrakardiālai lietošanai:

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Suns

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intravenozai lietošanai:

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Liellops

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Zirgs

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Suns

- Gaļa un blakusprodukti. no withdrawal period

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Anatomiski terapeitiski ķīmiskais veterinārais (ATĶvet) kods:

QN51AX50

Izplatīšanas juridiskie nosacījumi:

Recepšu veterinārās zāles

Atļaujas statuss:

Derīga

Atļautas:

Pieejamas tikai [Spanish](#) [Czech](#) [German](#) [Estonian](#) [English](#) [French](#) [Italian](#) [Dutch](#) [Portuguese](#) [Slovak](#) [Swedish](#) [Icelandic](#) [Norwegian](#)

Pieejams:

Belgium

Iepakojuma apraksts:

Pieejamas tikai [English](#)

Papildu informācija

Tiesību tips:

Marketing Authorisation

Veterināro zāļu atļaujas piešķiršanas juridiskais pamats:

Pieejamas tikai [English](#) [Italian](#)

Tirdzniecības atļaujas turētājs:

Intervet International B.V.

Tirdzniecības atļaujas datums:

5/11/1996

Ražošanas vietas sēriju izlaidei:

Intervet International GmbH

Atbildīgā iestāde:

Federal Agency For Medicines And Health Products

Atļaujas numurs:

BE-V179304

Atļaujas statusa maiņas datums:

13/04/2021

Lai uzzinātu par veterināro zāļu blakusparādībām, lūdzu, dodieties uz:
www.adrreports.eu/vet

Dokumenti

Zāļu apraksts

Šis dokuments neeksistē šādā valodā(latviešu). Jūs to varat atrast citā valodā zemāk.

Lietošanas instrukcija

Šis dokuments neeksistē šādā valodā(latviešu). Jūs to varat atrast citā valodā zemāk.

Marķējuma teksts

Šis dokuments neeksistē šādā valodā(latviešu). Jūs to varat atrast citā valodā zemāk.