

Exagon 400 mg/ml solution for injection

Atļautas

- Pentobarbital sodium

Zāļu identifikācija

Zāļu nosaukums:

Exagon 400 mg/ml solution for injection

Exagon vet. 400 mg/ml injekšjonsvāeske, oppløsning

Aktīvā viela:

Pieejamas tikai English

Mērķsugas:

Suns

Kaķis

Zaķis

Trusis

Jūrascūciņa

Kāmis

Pele

Žurka

Balodis

Liellops

Mājputni

Zirgs

Zirgs (ponijs)

Ūdele

Sesks
Cūka
Putni
Varde
Ķirzaka
Čūska
Ūdens bruņurupucis

Lietošanas veids:

intraperitoneālai lietošanai
intravenozai lietošanai
intrakardiālai lietošanai
Intrapulmonālai lietošanai

Sīkāka informācija par zālēm

Aktīvā viela un stiprums:

Pieejamas tikai English
400.00 miligrams(i) / 1.00 mililitrs(i)

Farmaceitiskā forma:

Šķīdums injekcijām

**Ierobežojumu periods pēc lietošanas veida:
intraperitoneālai lietošanai:**

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Zaķis

- Gaļa un blakusprodukti. no withdrawal period

Do not use in animals intended for human or animal consumption. Adequate measures must 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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Trusis

- Gaļa un blakusprodukti. no withdrawal period

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intravenozai lietošanai:

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Balodis

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- Ola. no withdrawal period

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Liellops

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- Piens. no withdrawal period

Do not use in animals intended for human or animal consumption. Adequate measures must 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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Mājputni

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Zirgs

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Zirgs (ponijs)

- Piens. no withdrawal period

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Cūka

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intrakardiālai lietošanai:

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Zaķis

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Cūka

- Gaļa un blakusprodukti. no withdrawal period

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Intrapulmonālai lietošanai:

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Balodis

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- Ola. no withdrawal period

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used for human or animal consumption.

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Mājputni

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Anatomiski terapeitiski ķīmiskais veterinārais (ATĶvet) kods:

QN51AA01

Izplatīšanas juridiskie nosacījumi:

Recepšu veterinārās zāles

Atļaujas statuss:

Derīga

Atļautas:

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Pieejams:

Norway

Iepakojuma apraksts:

Pieejamas tikai [English](#)

Pieejamas tikai [English](#)

Papildu informācija

Tiesību tips:

Marketing Authorisation

Veterināro zāļu atļaujas piešķiršanas juridiskais pamats:

Pieteikums ģenēriskām veterinārajām zālēm (Direktīvas 2001/82/EC 13.(1) pants)

Tirdzniecības atļaujas turētājs:

Vetviva Richter GmbH

Tirdzniecības atļaujas datums:

19/03/2014

Ražošanas vietas sēriju izlaidei:

Vetviva Richter GmbH

Atbildīgā iestāde:

Norwegian Medical Products Agency

Atļaujas numurs:

13-9536

Atļaujas statusa maiņas datums:

29/01/2019

Atsauces dalībvalsts:

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Procedūras numurs:

DE/V/0155/001

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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.

Visu dokumentu apvienotais fails

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.