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LIDOR 20 MG/ML SOLUTION FOR INJECTION FOR HORSES, DOGS AND CATS

Ima dovoljenje za promet

- Lidocaine hydrochloride monohydrate

Informacije o zdravilu

Ime zdravila:
LIDOR 20 MG/ML SOLUTION FOR INJECTION FOR HORSES, DOGS AND CATS

Učinkovina:
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 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:
Epiduralna uporaba
Subkutana uporaba

Perinevralna uporaba
Okularna uporaba
Intraartikularna uporaba

Podatki o zdravilu

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

Farmacevtska oblika:
Raztopina za injiciranje

Karenca glede na pot uporabe:
Epiduralna uporaba:

- **Horse**
  - Meat and offal. 3 day
  - Milk. 3 day

Subkutana uporaba:

- **Horse**
  - Meat and offal. 3 day
  - Milk. 3 day

Perinevralna uporaba:

- **Horse**
  - Meat and offal. 3 day
  - Milk. 3 day

Okularna uporaba:

- **Horse**

- Meat and offal. 3 day

- Milk. 3 day

Intraartikularna uporaba:

- **Horse**

  - Meat and offal. 3 day

  - Milk. 3 day

Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):
QN01BB02

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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Na voljo v:
Germany

Opis ovojnine zdravila:
Na voljo samo v English
Na voljo samo v English
Na voljo samo v English
Na voljo samo v English
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 Latvian Norwegian

Imetnik dovoljenja za promet z zdravilom:
Vetviva Richter GmbH

Datum dovoljenja za promet z zdravilom:
25/01/2018

Proizvodna mesta, odgovorna za sproščanje serij:
Vetviva Richter GmbH

Pristojni organ:
Federal Office Of Consumer Protection And Food Safety

Številka dovoljenja za promet z zdravilom:
402425.00.00

Datum spremembe statusa dovoljenja za promet:
25/01/2018

Referenčna država članica:
Na voljo samo v Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian

Številka postopka:
FR/V/0318/001

Zadevne države članice:
Na voljo samo v Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian
Na voljo samo v Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian
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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.

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