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Loxicom 20 mg/ml - Solution for injection (cattle, pigs, horses)

Ima dovoljenje za promet

- Meloxicam

Informacije o zdravilu

Ime zdravila:
Loxicom 20 mg/ml - Solution for injection (cattle, pigs, horses)

Učinkovina:
Na voljo samo v English

Ciljne živalske vrste:
govedo
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:
intramuskularna uporaba
Intravenska uporaba
Subkutana uporaba

Podatki o zdravilu

Učinkovina / Jakost:
Na voljo samo v English
Presentation_strength:20 mg Reference:B P Index:0

Farmacevtska oblika:
Raztopina za injiciranje

Karenca glede na pot uporabe:
intramuskularna uporaba:

- **govedo**
  - Meat and offal. 15 day 15 days
  - Milk. 5 day 5 days
- **Horse**
  - Meat and offal. 5 day 5 days
- **Pig**
  - Meat and offal. 5 day 5 days

Intravenska uporaba:

- **govedo**
  - Meat and offal. 15 day 15 days
  - Milk. 5 day 5 days
- **Horse**

- Meat and offal. 5 day 5 days

- **Pig**
  - Meat and offal. 5 day 5 days

Subkutana uporaba:

- **govedo**
  - Meat and offal. 15 day 15 days
  - Milk. 5 day 5 days

- **Horse**
  - Meat and offal. 5 day 5 days

- **Pig**
  - Meat and offal. 5 day 5 days

Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):
QM01AC06

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:
Belgium , Cyprus , Czechia , Denmark , Finland , Germany , Greece , Ireland , Italy

, Luxembourg , Netherlands , Poland , Portugal , Romania , Spain , Sweden , United Kingdom (Northern Ireland)

Opis ovojnine zdravila:
Na voljo samo v English
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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
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:
Norbrook Laboratories (Ireland) Limited

Datum dovoljenja za promet z zdravilom:
10/02/2009

Proizvodna mesta, odgovorna za sproščanje serij:
Norbrook Laboratories Limited
Norbrook Manufacturing Limited

Pristojni organ:
European Commission

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

Datum spremembe statusa dovoljenja za promet:
13/07/2011

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

Dokumenti

Combined File of all Documents

slovenščina (PDF)
Objavljeno na dan: 5/11/2026
Prenesi

ema-puar-loxicom-v-141-var-x-0003-en.pdf

ema-puar-loxicom-v-141-var-x-0005-en.pdf

ema-puar-loxicom-v-141-par-en.pdf

ema-puar-loxicom-v-141-var-x-0009-en.pdf