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TOLFENAMIC ACID VMD 40 mg/ml - Solution for injection

Ima dovoljenje za promet

- Tolfenamic acid

Informacije o zdravilu

Ime zdravila:
TOLFENAMIC ACID VMD 40 mg/ml - Solution for injection

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
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
40.00 milligram(s) / 1.00 millilitre(s)

Farmacevtska oblika:

Raztopina za injiciranje

Karenca glede na pot uporabe:

intramuskularna uporaba:

- **govedo**
  - Meat and offal. 12 day 12 days
  - Milk. 0 hour 0 hours
  - Meat and offal. 4 day 4 days
  - Milk. 24 hour 24 hours
- **Pig**
  - Meat and offal. 16 day 16 days

Intravenska uporaba:

- **govedo**
  - Meat and offal. 12 day 12 days
  - Milk. 0 hour 0 hours
  - Meat and offal. 4 day 4 days
  - Milk. 24 hour 24 hours

- **Pig**
  - Meat and offal. 16 day 16 days

Subkutana uporaba:

- **govedo**
  - Meat and offal. 12 day 12 days
  - Milk. 0 hour 0 hours
  - Meat and offal. 4 day 4 days
  - Milk. 24 hour 24 hours
- **Pig**
  - Meat and offal. 16 day 16 days

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

QM01AG02

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 , Czechia , France , Luxembourg , Netherlands , Poland , Slovakia , Slovenia

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Opis ovojnine zdravila:

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 French Italian Latvian Lithuanian Norwegian

Imetnik dovoljenja za promet z zdravilom:
V.M.D.

Datum dovoljenja za promet z zdravilom:
3/02/2025

Proizvodna mesta, odgovorna za sproščanje serij:
V.M.D.
Laboratoires Biove

Pristojni organ:
European Commission

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

Datum spremembe statusa dovoljenja za promet:
3/02/2025

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: 13/03/2026
Prenesi

ema-puar-v6234-tolfenamicacidvmd-initial-en.pdf