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Metacam 5 mg/ml - Solution for injection (cattle, pigs)

Ima dovoljenje za promet

- Meloxicam

Informacije o zdravilu

Ime zdravila:
Metacam 5 mg/ml - Solution for injection (cattle, pigs)

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

Pot uporabe:
intramuskularna uporaba
Intravenska uporaba
Subkutana uporaba

Podatki o zdravilu

Učinkovina / Jakost:
Na voljo samo v English

Presentation_strength:5 mg Reference:BP Index:0

Farmacevtska oblika:
Raztopina za injiciranje

Karenca glede na pot uporabe:
intramuskularna uporaba:

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

Intravenska uporaba:

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

Subkutana uporaba:

- **govedo**
  - Meat and offal. 15 day 15 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 samo v Estonian English French Lithuanian Portuguese Swedish Icelandic Norwegian

Na voljo v:
Austria , Belgium , Czechia , Luxembourg , Poland

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
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

Imetnik dovoljenja za promet z zdravilom:
Boehringer Ingelheim Vetmedica GmbH

Datum dovoljenja za promet z zdravilom:
7/01/1998

Proizvodna mesta, odgovorna za sproščanje serij:

Labiana Life Sciences S.A.

Pristojni organ:
European Commission

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

Datum spremembe statusa dovoljenja za promet:
7/01/1998

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: 14/03/2026
Prenesi

ema-puar-metacam-v-033-var-x-0119-en.pdf

ema-puar-metacam-v-033-var-x-0107-en.pdf

ema-puar-metacam-v-033-par-en.pdf

ema-puar-metacam-novem-var-ws-0264-en.pdf

ema-puar-metacam-v-033-var-ii-0127-en.pdf

ema-puar-v33-metacam-ii0084-en.pdf

ema-puar-v33-metacam-vra0151g-en.pdf