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# Labiprofen 150 mg/ml raztopina za injiciranje za govedo, prašiče in konje

Ima dovoljenje za promet

- Ketoprofen

## Informacije o zdravilu

**Ime zdravila:**
Labiprofen 150 mg/ml raztopina za injiciranje za govedo, prašiče in konje

**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
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

# Podatki o zdravilu

**Učinkovina / Jakost:**

Na voljo samo v English
150.00 milligram(s) / 1.00 millilitre(s)

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**Farmacevtska oblika:**

Raztopina za injiciranje

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**Karenca glede na pot uporabe:**

**intramuskularna uporaba:**

- **Pig**
  - Meat and offal. 3 day
- **govedo**
  - Meat and offal. 2 day
  - Milk. 0 hour

**Intravenska uporaba:**

- **Horse**
  - Meat and offal. 1 day
  - Milk. no withdrawal period

Milk: Not authorised for use in mares producing milk for human consumption.

- **govedo**
  - Meat and offal. 2 day
  - Milk. 0 hour

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

QM01AE03

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**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

**Na voljo v:**
Slovenia

**Opis ovojnine zdravila:**
Kartonasta škatla z 1 vialo po 50 ml
Kartonasta škatla z 1 vialo po 100 ml
Kartonasta škatla z 1 vialo po 250 ml
Kartonasta škatla z 12 vialami po 50 ml
Kartonasta škatla z 10 vialami po 100 ml
Kartonasta škatla z 10 vialami po 250 ml

## 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:**
Labiana Life Sciences S.A.

**Datum dovoljenja za promet z zdravilom:**
31/03/2021

**Proizvodna mesta, odgovorna za sproščanje serij:**
Labiana Life Sciences S.A.

**Pristojni organ:**
Agency For Medicinal Products And Medical Devices Of The Republic Of Slovenia

**Številka dovoljenja za promet z zdravilom:**
DC/V/0731/001

**Datum spremembe statusa dovoljenja za promet:**
31/03/2021

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

**Številka postopka:**
ES/V/0388/001

**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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Na voljo samo v Estonian English French Lithuanian Portuguese 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

Povzetek glavnih značilnosti zdravila

slovenščina (PDF)
Objavljeno na dan: 17/04/2026
Prenesi

Navodilo za uporabo

slovenščina (PDF)
Objavljeno na dan: 17/04/2026
Prenesi

Označevanje

slovenščina (PDF)
Objavljeno na dan: 17/04/2026
Prenesi

Combined File of all Documents

Ta dokument ne obstaja v tem jeziku (slovenščina). Lahko ga najdete spodaj v drugem jeziku.