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EPRINOVET MULTI 5 MG/ML POUR-ON SOLUTION FOR CATTLE, SHEEP AND GOATS

Ima dovoljenje za promet

- Eprinomectin

Informacije o zdravilu

Ime zdravila:
EPRINOVET MULTI 5 MG/ML POUR-ON SOLUTION FOR CATTLE, SHEEP AND GOATS

Učinkovina:
Na voljo samo v English

Ciljne živalske vrste:
pitovno govedo
Na voljo samo v Spanish Czech Danish Estonian Greek English French Italian Latvian Lithuanian Romanian Finnish Swedish 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:
Dajanje kožnega poliva

Podatki o zdravilu

Učinkovina / Jakost:

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

Farmacevtska oblika:

Kožni poliv, raztopina

Karenca glede na pot uporabe:

Dajanje kožnega poliva:

- **pitovno govedo**
  - Meat and offal. 15 day
- **Cattle (dairy cow)**
  - Milk. 0 hour
- **Sheep**
  - Meat and offal. 2 day
- **Sheep**
  - Milk. 0 hour
- **Goat**
  - Meat and offal. 1 day
- **Goat**
  - Milk. 0 hour

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

QP54AA04

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

Opis ovojnine zdravila:
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:
Laboratorios Calier S.A.

Datum dovoljenja za promet z zdravilom:
11/08/2026

Proizvodna mesta, odgovorna za sproščanje serij:
Laboratorios Calier S.A.

Pristojni organ:
Institute For State Control Of Veterinary Biologicals And Medicaments

Številka dovoljenja za promet z zdravilom:
96/032/DC/26-S

Datum spremembe statusa dovoljenja za promet:
11/08/2026

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/0443/001

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

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.

Povzetek glavnih značilnosti zdravila

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

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