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Equiparin 50.000 IU/100 g Gel for Horses

Ima dovoljenje za promet

- Levomenthol
- Hydroxyethyl salicylate
- Heparin sodium

Informacije o zdravilu

Ime zdravila:
Equiparin 50.000 IU/100 g Gel for Horses

Učinkovina:
Na voljo samo v English
Na voljo samo v English
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

Pot uporabe:
Dermalna uporaba

Podatki o zdravilu

Učinkovina / Jakost:

Na voljo samo v English
0.50 gram(s) / 100.00 gram(s)

Na voljo samo v English
5.00 gram(s) / 100.00 gram(s)

Na voljo samo v English
50000.00 international unit(s) / 100.00 gram(s)

Farmacevtska oblika:

Gel

Karenca glede na pot uporabe:

Dermalna uporaba:

- **Horse**
  - Meat and offal. 0 day
  - Milk. no withdrawal period

Not permitted for use in mares producing milk for human consumption.

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

QM02AC99

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 Estonian English French Lithuanian Portuguese Swedish Icelandic Norwegian

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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 Italian Latvian Norwegian

Imetnik dovoljenja za promet z zdravilom:
aniMedica GmbH

Datum dovoljenja za promet z zdravilom:
29/01/2010

Proizvodna mesta, odgovorna za sproščanje serij:
aniMedica GmbH

Pristojni organ:
The Veterinary Medicines Directorate

Številka dovoljenja za promet z zdravilom:
Vm 24745/4004

Datum spremembe statusa dovoljenja za promet:
12/12/2014

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

Številka postopka:
DE/V/0128/002

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

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