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Equibactin vet. 250 mg/g + 50 mg/g, oral powder

Ima dovoljenje za promet

- Trimethoprim
- Sulfadiazine

Informacije o zdravilu

Ime zdravila:
Equibactin vet. 250 mg/g + 50 mg/g, oral powder

Učinkovina:
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:
Peroralna uporaba

Podatki o zdravilu

Učinkovina / Jakost:
Na voljo samo v English
50.00 milligram(s) / 1.00 gram(s)

Na voljo samo v English
250.00 milligram(s) / 1.00 gram(s)

Farmacevtska oblika:

Peroralni prašek

Karenca glede na pot uporabe:

Peroralna uporaba:

- **Horse**
  - Meat and offal. 20 day

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

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

QJ01EW10

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:

Norway

Opis ovojnine zdravila:

Na voljo samo v Swedish
Na voljo samo v Swedish
Na voljo samo v Swedish
Na voljo samo v Swedish
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Na voljo samo v Swedish

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:
Dechra Regulatory B.V.

Datum dovoljenja za promet z zdravilom:
27/05/2019

Proizvodna mesta, odgovorna za sproščanje serij:
Lelypharma B.V.

Pristojni organ:
Norwegian Medical Products Agency

Številka dovoljenja za promet z zdravilom:
17-11685

Datum spremembe statusa dovoljenja za promet:
27/05/2019

Referenčna država članica:

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

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Številka postopka:
SE/V/0120/001

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

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

Navodilo za uporabo

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

Označevanje

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