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Triquest Vet, 333 mg/ml + 67 mg/ml, oral suspension for horses

Ima dovoljenje za promet

- Sulfadiazine
- Trimethoprim

Informacije o zdravilu

Ime zdravila:
Triquest Vet, 333 mg/ml + 67 mg/ml, oral suspension for horses

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
Peroralna uporaba

Podatki o zdravilu

Učinkovina / Jakost:

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

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

Farmacevtska oblika:

Peroralna suspenzija

Karenca glede na pot uporabe:

Peroralna uporaba:

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

Peroralna uporaba:

- **Horse**
  - Milk. no withdrawal period

Milk: Not authorised for use in animals 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

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Opis ovojnine zdravila:
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 French Italian Latvian Lithuanian Norwegian

Imetnik dovoljenja za promet z zdravilom:
Alfasan Nederland B.V.

Datum dovoljenja za promet z zdravilom:
3/10/2024

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

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

Številka dovoljenja za promet z zdravilom:
DC/V/0812/001

Datum spremembe statusa dovoljenja za promet:
3/10/2024

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

Številka postopka:
SE/V/0125/001

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

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

Combined File of all Documents

slovenščina (PDF)
Objavljeno na dan: 4/11/2026
Prenesi

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.

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