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Forespix, 100mg/ml, Solution for injection

Ima dovoljenje za promet

- Tulathromycin

Informacije o zdravilu

Ime zdravila:
Forespix, 100mg/ml, Solution for injection

Učinkovina:
Na voljo samo v English

Ciljne živalske vrste:
govedo
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:
Subkutana uporaba
intramuskularna uporaba

Podatki o zdravilu

Učinkovina / Jakost:
Na voljo samo v English
100.00 milligram(s) / 1.00 millilitre(s)

Farmacevtska oblika:
Raztopina za injiciranje

Karenca glede na pot uporabe:
Subkutana uporaba:

- **govedo**
  - Meat and offal. 22 day
  - Milk. no withdrawal period

Do not use in pregnant animals intended for milk production for human consumption during 2 months prior to expected parturition.,

intramuskularna uporaba:

- **Pig**
  - Meat and offal. 13 day
- **Sheep**
  - Meat and offal. 16 day
  - Milk. no withdrawal period

Do not use in pregnant animals intended for milk production for human consumption during 2 months prior to expected parturition.,

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Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):
QJ01FA94

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 Latvian Dutch Portuguese Slovak Swedish Icelandic Norwegian

Na voljo v:
Czechia

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 Italian Latvian Norwegian

Imetnik dovoljenja za promet z zdravilom:
Przedsiebiorstwo Wielobranzowe Vet-Agro Sp. z o.o.

Datum dovoljenja za promet z zdravilom:
16/12/2020

Proizvodna mesta, odgovorna za sproščanje serij:
Przedsiebiorstwo Wielobranzowe Vet-Agro Sp. z o.o.
Przedsiebiorstwo Wielobranzowe Vet-Agro Sp. z o.o.

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

Številka dovoljenja za promet z zdravilom:
96/077/20-C

Datum spremembe statusa dovoljenja za promet:
16/12/2020

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

Številka postopka:
CZ/V/0166/001

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 Croatian Italian Dutch Portuguese Slovak 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.

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