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Taneven 300 mg/ml suspension for injection for horses, cattle, sheep, goats, dogs and cats

Ima dovoljenje za promet

- Benzylpenicillin (procaine) monohydrate

Informacije o zdravilu

Ime zdravila:
Taneven 300 mg/ml suspension for injection for horses, cattle, sheep, goats, dogs and cats

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
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
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
300.00 milligram(s) / 1.00 millilitre(s)

Farmacevtska oblika:
Suspenzija za injiciranje

Karenca glede na pot uporabe:
Subkutana uporaba:

- **govedo**
  - Meat and offal. 10 day
  - Milk. 120 hour
- **Goat**
  - Milk. 120 hour
  - Meat and offal. 10 day
- **Sheep**
  - Meat and offal. 10 day
  - Milk. 120 hour
-

Horse

- Meat and offal. 10 day
- Milk. no withdrawal period

Not authorised for use mares producing milk for human consumption.

intramuskularna uporaba:

- **govedo**
  - Meat and offal. 10 day
  - Milk. 120 hour
- **Goat**
  - Milk. 120 hour
  - Meat and offal. 10 day
- **Sheep**
  - Meat and offal. 10 day
  - Milk. 120 hour
- **Horse**
  - Meat and offal. 10 day
  - Milk. no withdrawal period

Not authorised for use mares producing milk for human consumption.

Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):
QJ01CE09

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

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:
Wirtschaftsgenossenschaft deutscher Tieraerzte eG

Datum dovoljenja za promet z zdravilom:
15/12/2021

Proizvodna mesta, odgovorna za sproščanje serij:
Wirtschaftsgenossenschaft deutscher Tieraerzte eG

Pristojni organ:
Ministry Of Health

Številka dovoljenja za promet z zdravilom:
105454

Datum spremembe statusa dovoljenja za promet:
15/12/2021

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

Številka postopka:
DE/V/0337/001

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 Spanish Czech German Estonian English French Italian Lithuanian Dutch Portuguese Romanian 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.

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