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Taneven LC 3 g intramammary suspension for lactating cows

Ima dovoljenje za promet

- Benzylpenicillin (procaine) monohydrate

Informacije o zdravilu

Ime zdravila:
Taneven LC 3 g intramammary suspension for lactating cows

Učinkovina:
Na voljo samo v English

Ciljne živalske vrste:
govedo

Pot uporabe:
Intramamarno dajanje

Podatki o zdravilu

Učinkovina / Jakost:
Na voljo samo v English
3.00 gram(s) / 1.00 Aplikator

Farmacevtska oblika:
intramamarna suspenzija

Karenca glede na pot uporabe:

Intramamarno dajanje:

- **govedo**
  - Milk. 120 hour
  - Meat and offal. 5 day

Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):
QJ51CE09

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
Na voljo samo v English
Na voljo samo v English
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

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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:
30/11/2022

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

Pristojni organ:
Ministere De La Sante Division De La Pharmacie Et Des Medicaments

Številka dovoljenja za promet z zdravilom:
V 110/22/11/0001

Datum spremembe statusa dovoljenja za promet:
30/11/2022

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

Številka postopka:
DE/V/0338/001

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