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Utertab 2000 mg intrauterine tablet for cattle

Ima dovoljenje za promet

- Tetracycline hydrochloride

Informacije o zdravilu

Ime zdravila:
Utertab 2000 mg intrauterine tablet for cattle

Učinkovina:
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:
Intrauterina uporaba

Podatki o zdravilu

Učinkovina / Jakost:
Na voljo samo v English
2000.00 milligram(s) / 1.00 Tableta

Farmacevtska oblika:

intrauterina tableta

Karenca glede na pot uporabe:

Intrauterina uporaba:

- **Cattle (lactating cow)**
  - Milk. 96 hour
  - Meat and offal. 10 day

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

QG51AA02

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:

Germany

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

Pravna podlaga za izdajo dovoljenja za promet z zdravilom:
Na voljo samo v English Italian Latvian Norwegian

Imetnik dovoljenja za promet z zdravilom:
aniMedica GmbH

Datum dovoljenja za promet z zdravilom:
20/08/2018

Proizvodna mesta, odgovorna za sproščanje serij:
aniMedica GmbH
aniMedica Herstellungs GmbH

Pristojni organ:
Federal Office Of Consumer Protection And Food Safety

Številka dovoljenja za promet z zdravilom:
402466.00.00

Datum spremembe statusa dovoljenja za promet:
20/08/2018

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

Številka postopka:
DE/V/0176/001

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 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

Combined File of all Documents

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

2402466-paren-20181009.rtf