Oxytetraciclin hydrochloride-NGP 7/ 0,340 g, 11/ 0,510 g, 22/ 1,020 g comprettae spumescentes

Volitatud

- Oxytetracycline hydrochloride
- Oxytetracycline hydrochloride
- Oxytetracycline hydrochloride

Product identification

Ravimi nimetus:
Окситетрациклин хидрохлорид-NGP 7/ 0,340 g, 11/ 0,510 g, 22/ 1,020 g пенообразуващи компрети
Oxytetraciclin hydrochloride-NGP 7/ 0,340 g, 11/ 0,510 g, 22/ 1,020 g comprettae spumescentes

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Toimeaine:
Turustatakse ainult English
Turustatakse ainult English
Turustatakse ainult English

Loomaliigid:
siga (suguemis)
kits
lammas

Manustamisviis:
Intrauteriinne

Product details

Toimeaine / Tugevus:

Turustatakse ainult English
0.34 gram(s) / 1.00 Tablett

Turustatakse ainult English
0.51 gram(s) / 1.00 Tablett

Turustatakse ainult English
1.02 gram(s) / 1.00 Tablett

Ravimvorm:

Intrauteriintablett

Withdrawal period by route of administration:

Intrauteriinne:

- **siga (suguemis)**

  - liha ja söödavad koed. no withdrawal period

Месо и вътрешни органи - третираните животни не може да се използват за консумация от хора

- **kits**

  - piim. no withdrawal period

Мляко - по време на третирането и най-малко 4-5 дни след това животните отделят коластра, която не се консумира от хора

  - liha ja söödavad koed. no withdrawal period

Месо и вътрешни органи - третираните животни не може да се използват за консумация от хора

- **lammas**

- liha ja söödavad koed. no withdrawal period

Месо и вътрешни органи - третираните животни не може да се използват за консумация от хора

- piim. no withdrawal period

Мляко - по време на третирането и най-малко 4-5 дни след това животните отделят коластра, която не се консумира от хора

Veterinaarravimite anatoomilis-terapeutilise keemilise klassifikatsiooni (ATCvet) kohane kood:
QJ01AA

Õiguslik tarnestaatus:
Retseptiravim - veterinaarravim

Müügiloa staatus:
Valid

Authorised in:
Bulgaaria

Pakendi kirjeldus:
Turustatakse ainult Bulgarian
Turustatakse ainult Bulgarian
Turustatakse ainult Bulgarian
Turustatakse ainult Bulgarian

Additional information

Entitlement type:
Turustatakse ainult English French Croatian Italian Latvian Finnish Swedish Icelandic Norwegian

Ravimi müügiloa õiguslik alus:
Turustatakse ainult English Italian Latvian Norwegian

Müügiloa hoidja:

Northern Veterinary Dealer-SVD OOD

Marketing authorisation date:
10/06/2007

Partii vabastamise tootmiskohad:
Northern Veterinary Dealer-SVD OOD

Vastutav asutus:
Bulgarian Food Safety Authority

Authorisation number:
0022-1649-03-11-2011

Müügiloa staatuse muutmise kuupäev:
2/11/2011

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Ravimi omaduste kokkuvõte

Seda dokumenti selles keeles ei eksisteeri (eesti). Selle leiate allpool teises keeles.

Package Leaflet and Labelling

Seda dokumenti selles keeles ei eksisteeri (eesti). Selle leiate allpool teises keeles.
