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Oxtra 10%,92.7mg/ml, ενέσιμο διάλυμα για βοοειδή, άλογα, χοίρους και πρόβατα

Ima dovoljenje za promet

- Oxytetracycline hydrochloride

Informacije o zdravilu

Ime zdravila:
Oxtra 10%,92.7mg/ml, ενέσιμο διάλυμα για βοοειδή, άλογα, χοίρους και πρόβατα

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

Pot uporabe:
intramuskularna uporaba

Intravenska 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:

intramuskularna uporaba:

- **govedo**
  - Meat and offal. 15 day
  - Milk. 144 hour
- **Sheep**
  - Meat and offal. 20 day
  - Milk. 216 hour
- **Horse**
  - Meat and offal. 23 day
  - Milk. no withdrawal period

NOT AUTHORIZED FOR USE IN LACTATING ANIMALS PRODUCING MILK FOR HUMAN CONSUPMTION

- **Pig**
  - Meat and offal. 8 day

Intravenska uporaba:

-

govedo

- Milk. 144 hour
- Meat and offal. 15 day

•

Sheep

- Meat and offal. 20 day
- Milk. 216 hour

•

Horse

- Meat and offal. 23 day
- Milk. no withdrawal period

NOT AUTHORIZED FOR USE IN LACTATING ANIMALS PRODUCING MILK FOR HUMAN CONSUPMTION

•

Pig

- Meat and offal. 8 day

Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):
QJ01AA06

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 Greek English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian

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Na voljo v:
Cyprus

Opis ovojnine zdravila:

Na voljo samo v Greek

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 Lithuanian Norwegian

Imetnik dovoljenja za promet z zdravilom:
Fatro S.p.A.

Datum dovoljenja za promet z zdravilom:
7/05/1986

Proizvodna mesta, odgovorna za sproščanje serij:
Fatro S.p.A.

Pristojni organ:
Veterinary Services, Ministry Of Agriculture, Natural Resources And Environment

Številka dovoljenja za promet z zdravilom:
10460

Datum spremembe statusa dovoljenja za promet:
30/05/2012

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.