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Receptal 4 microgram/ml solution for injection

Ima dovoljenje za promet

- Buserelin

Informacije o zdravilu

Ime zdravila:
Receptal 4 microgram/ml solution for injection

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 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 Estonian English French Italian Latvian Lithuanian Hungarian Dutch Romanian Finnish Swedish Norwegian

Pot uporabe:
intramuskularna uporaba

Intravenska uporaba
Subkutana uporaba

Podatki o zdravilu

Učinkovina / Jakost:
Na voljo samo v English
4.00 microgram(s) / 1.00 millilitre(s)

Farmacevtska oblika:
Raztopina za injiciranje

Karenca glede na pot uporabe:
intramuskularna uporaba:

- **govedo**
  - Meat and offal. 0 day
  - Milk. 0 day
- **Horse**
  - Meat and offal. 0 day
- **Rabbit**
  - Meat and offal. 0 day
- **Pig**
  - Meat and offal. 0 day
- **Trout**
  - Meat and offal. 0 day

Intravenska uporaba:

-

govedo

- Meat and offal. 0 day
- Milk. 0 day

•

Horse

- Meat and offal. 0 day

•

Rabbit

- Meat and offal. 0 day

•

Pig

- Meat and offal. 0 day

Subkutana uporaba:

•

govedo

- Meat and offal. 0 day
- Milk. 0 day

•

Horse

- Meat and offal. 0 day

•

Rabbit

- Meat and offal. 0 day

•

Pig

- Meat and offal. 0 day

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

QH01CA90

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
Na voljo samo v English
Na voljo samo v English
Na voljo samo v English
Na voljo samo v English
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Na voljo samo v English
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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

Imetnik dovoljenja za promet z zdravilom:
Intervet Nederland B.V.

Datum dovoljenja za promet z zdravilom:
8/09/1992

Proizvodna mesta, odgovorna za sproščanje serij:
Intervet International GmbH

Pristojni organ:
Medicines Evaluation Board

Številka dovoljenja za promet z zdravilom:
REG NL 5327

Datum spremembe statusa dovoljenja za promet:
25/01/2017

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.