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Paramore, 200 mg/ml, soluzione per uso orale per vitelli da latte, suini, polli, tacchini e conigli

Ima dovoljenje za promet

- Paromomycin sulfate

Informacije o zdravilu

Ime zdravila:
Paramore, 200 mg/ml, soluzione per uso orale per vitelli da latte, suini, polli, tacchini e conigli

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
Na voljo samo v Bulgarian Spanish Czech Danish German Estonian Greek English French Italian Latvian Lithuanian Hungarian Dutch Romanian Finnish Swedish Icelandic Norwegian
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Na voljo samo v Bulgarian Spanish Danish Estonian English French Italian Latvian Lithuanian Hungarian Dutch Romanian Finnish Swedish Norwegian

Pot uporabe:
Peroralna uporaba

Podatki o zdravilu

Učinkovina / Jakost:
Na voljo samo v English
200.00 milligram(s) / 1.00 millilitre(s)

Farmacevtska oblika:
Peroralna raztopina

Karenca glede na pot uporabe:
Peroralna uporaba:

- **Turkey**
  - Meat and offal. 2 day

Uso non autorizzato in tacchini che producono uova per il consumo umano

- **Chicken**
  - Meat and offal. 2 day
  - Eggs. 0 day
- **Rabbit**
  - Meat and offal. 3 day
- **Pig**

- Meat and offal. 44 day

- **Cattle (suckling calf)**
  - Meat and offal. 44 day

Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):
QA07AA06

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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Na voljo v:
Italy

Opis ovojnine zdravila:
Na voljo samo v Italian
Na voljo samo v Italian

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:
HuVepharma

Datum dovoljenja za promet z zdravilom:
11/04/2016

Proizvodna mesta, odgovorna za sproščanje serij:
Biovet AD

Pristojni organ:
Ministry Of Health

Številka dovoljenja za promet z zdravilom:
Ta podatek za to zdravilo ni na voljo.

Datum spremembe statusa dovoljenja za promet:
11/04/2016

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.