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PARACOX-5, SUSPENSION FOR ORAL SUSPENSION FOR CHICKENS

Ima dovoljenje za promet

- Eimeria maxima, strain CP, Live
- Eimeria tenella, strain HP, Live
- Eimeria mitis, strain HP, Live
- Eimeria maxima, strain MFP, Live
- Eimeria acervulina, strain HP, Live

Informacije o zdravilu

Ime zdravila:
PARACOX-5, SUSPENSION FOR ORAL SUSPENSION FOR CHICKENS

Učinkovina:
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

Ciljne živalske vrste:
brojler
Na voljo samo v Spanish Danish Estonian English French Italian Latvian Lithuanian Romanian Finnish Swedish Icelandic Norwegian

Pot uporabe:

Peroralna uporaba

Podatki o zdravilu

Učinkovina / Jakost:

Na voljo samo v English
500.00 Organisms / 0.00 millilitre(s)

Na voljo samo v English
1000.00 Organisms / 0.00 millilitre(s)

Na voljo samo v English
100.00 Organisms / 0.00 millilitre(s)

Na voljo samo v English
200.00 Organisms / 0.00 millilitre(s)

Na voljo samo v English
500.00 Organisms / 0.00 millilitre(s)

Farmacevtska oblika:

Peroralna suspenzija

Karenca glede na pot uporabe:

Peroralna uporaba:

- **brojler**
  - All relevant tissues. 0 day
- **Chicken (one day-old chick)**
  - All relevant tissues. 0 day

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

QI01AN01

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

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 French Italian Latvian Norwegian

Imetnik dovoljenja za promet z zdravilom:
Intervet Hellas M.A.E.

Datum dovoljenja za promet z zdravilom:
27/06/1999

Proizvodna mesta, odgovorna za sproščanje serij:
Merck Sharp & Dohme Animal Health S.L.
MSD Animal Health UK Limited

Pristojni organ:
National Organization For Medicines

Številka dovoljenja za promet z zdravilom:
69553/22-11-2005/K-0136701

Datum spremembe statusa dovoljenja za promet:
15/07/2021

Referenčna država članica:

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

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Številka postopka:

FR/V/0351/001

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 Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian

Na voljo samo v Spanish Czech German Estonian Greek English French 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.