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HuveGuard MMAT suspension for oral suspension for chickens

Ima dovoljenje za promet

- Eimeria tenella, strain Rt 3+15, Live
- Eimeria acervulina, strain RA 3+20, Live
- Eimeria mitis, strain Jormit 3+9, Live
- Eimeria maxima, strain MCK +10, Live

Informacije o zdravilu

Ime zdravila:
HuveGuard MMAT 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

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

Pot uporabe:
Peroralna uporaba

Podatki o zdravilu

Učinkovina / Jakost:

Na voljo samo v English
417.00 oocyst(s) / 0.03 millilitre(s)

Na voljo samo v English
50.00 unit(s) / 0.03 millilitre(s)

Na voljo samo v English
278.00 unit(s) / 0.03 millilitre(s)

Na voljo samo v English
278.00 unit(s) / 0.03 millilitre(s)

Farmacevtska oblika:

Suspenzija za peroralno suspenzijo

Karenca glede na pot uporabe:

Peroralna uporaba:

- **Chicken**
  - Meat and offal. 0 day
  - Egg. 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 German Estonian English French Italian Dutch Portuguese Slovak Finnish Swedish Icelandic Norwegian

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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
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 Latvian Lithuanian Norwegian

Imetnik dovoljenja za promet z zdravilom:
HuVepharma

Datum dovoljenja za promet z zdravilom:
6/07/2016

Proizvodna mesta, odgovorna za sproščanje serij:
Biovet AD

Pristojni organ:
State Agency Of Medicines

Številka dovoljenja za promet z zdravilom:
1985

Datum spremembe statusa dovoljenja za promet:
6/07/2016

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

Številka postopka:

NL/V/0206/001

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

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Na voljo samo v Estonian English French Lithuanian Portuguese 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

Povzetek glavnih značilnosti zdravila

Ta dokument ne obstaja v tem jeziku (slovenščina). Lahko ga najdete spodaj v drugem jeziku.

Combined File of all Documents

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