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AviPro THYMOVAC Lyophilisate for use in drinking water

Ima dovoljenje za promet

- Chicken anaemia virus, strain CUX-1, Live

Informacije o zdravilu

Ime zdravila:
AviPro THYMOVAC Lyophilisate for use in drinking water

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

Pot uporabe:
Dajanje v vodo za pitje

Podatki o zdravilu

Učinkovina / Jakost:
Na voljo samo v English
31622.00 50% tissue culture infectious dose / 1.00 Dose

Farmacevtska oblika:
Liofilizat za dajanje v vodo za pitje

Karenca glede na pot uporabe:
Dajanje v vodo za pitje:

- **Chicken**
  - Egg. 0 day
  - Meat and offal. 0 day

Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):
QI01AD04

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

Na voljo v:
Sweden

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
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 Norwegian

Imetnik dovoljenja za promet z zdravilom:
Lohmann Animal Health GmbH

Datum dovoljenja za promet z zdravilom:
4/12/2009

Proizvodna mesta, odgovorna za sproščanje serij:
Lohmann Animal Health GmbH

Pristojni organ:
Swedish Medical Products Agency

Številka dovoljenja za promet z zdravilom:
26904

Datum spremembe statusa dovoljenja za promet:
4/12/2009

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

Številka postopka:
DE/V/0247/001

Zadevne države članice:

Na voljo samo v Spanish Czech German Estonian 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

Navodilo za uporabo

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

Povzetek glavnih značilnosti zdravila

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

Označevanje

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Combined File of all Documents

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