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AviPro IBD Xtreme Lyophilisate for use in drinking water for chickens

Ima dovoljenje za promet

- Infectious bursal disease virus, strain V217 (intermediate plus), Live

Informacije o zdravilu

Ime zdravila:
AviPro IBD Xtreme Lyophilisate for use in drinking water for chickens

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

31.00 50% Embryo Infective Dose / 1.00 Dose

Farmacevtska oblika:
Liofilizat za dajanje v vodo za pitje

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

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

Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):
QI01AD09

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:
Germany

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:
27/02/2008

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

Pristojni organ:
Paul-Ehrlich-Institut

Številka dovoljenja za promet z zdravilom:
PEI.V.03418.01.1

Datum spremembe statusa dovoljenja za promet:
28/11/2012

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

Številka postopka:
DE/V/0238/001

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 Greek 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

Combined File of all Documents

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