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Avishield IB H120, lyophilisate for oculonasal suspension/use in drinking water, for chickens

Ima dovoljenje za promet

- Infectious bronchitis virus, type Massachusetts, strain H120, Live

Informacije o zdravilu

Ime zdravila:
Avishield IB H120, lyophilisate for oculonasal suspension/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:
Okulonazalno dajanje
Dajanje v vodo za pitje

Podatki o zdravilu

Učinkovina / Jakost:

Na voljo samo v English
3.50 log 10 50% embryo infective dose / 1.00 Dose

Farmacevtska oblika:

Liofilizat za okulonazalno suspenzijo/dajanje v vodo za pitje

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Karenca glede na pot uporabe:

Okulonazalno dajanje:

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

Dajanje v vodo za pitje:

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

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

QI01AD07

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 Latvian Dutch Portuguese Slovak 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

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:
Izo S.r.l.

Datum dovoljenja za promet z zdravilom:
23/01/2018

Proizvodna mesta, odgovorna za sproščanje serij:
Genera d.d.

Pristojni organ:
Institute For State Control Of Veterinary Biologicals And Medicaments

Številka dovoljenja za promet z zdravilom:
97/005/18-C

Datum spremembe statusa dovoljenja za promet:
28/01/2022

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

Številka postopka:
NL/V/0292/001

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

Combined File of all Documents

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

Povzetek glavnih značilnosti zdravila

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Navodilo za uporabo

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Označevanje

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