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AVINEW

Ima dovoljenje za promet

- Newcastle disease virus, strain VG/GA-AVINEW, Live

Informacije o zdravilu

Ime zdravila:
AVINEW

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
Na voljo samo v Bulgarian Spanish Czech Danish German Estonian Greek English French Italian Latvian Lithuanian Hungarian Dutch Romanian Finnish Swedish Icelandic Norwegian
brojler

Pot uporabe:
Okularna uporaba
Peroralna uporaba
Okulonazalno dajanje

Podatki o zdravilu

Učinkovina / Jakost:

Na voljo samo v English
5.50 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:

Okularna uporaba:

- **Chicken (layer hen)**
  - All relevant tissues. 0 day
- **Chicken (for reproduction)**
  - All relevant tissues. 0 day
- **brojler**
  - All relevant tissues. 0 day

Peroralna uporaba:

- **Chicken (layer hen)**
  - All relevant tissues. 0 day
- **Chicken (for reproduction)**
  - All relevant tissues. 0 day
- **brojler**
  - All relevant tissues. 0 day

Okulonazalno dajanje:

- **Chicken (layer hen)**
  - All relevant tissues. 0 day
- **Chicken (for reproduction)**
  - All relevant tissues. 0 day
- **brojler**
  - All relevant tissues. 0 day

Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):
QI01AD06

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

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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

Dodatne informacije

Vrsta dovoljenja:
Na voljo samo v English French Croatian Italian Latvian Finnish Swedish Icelandic Norwegian

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Pravna podlaga za izdajo dovoljenja za promet z zdravilom:
Na voljo samo v English French Italian Latvian Norwegian

Imetnik dovoljenja za promet z zdravilom:
Boehringer Ingelheim Animal Health France

Datum dovoljenja za promet z zdravilom:
13/12/1999

Proizvodna mesta, odgovorna za sproščanje serij:
Boehringer Ingelheim Animal Health France

Pristojni organ:
French Agency For Food, Environmental And Occupational Health & Safety

Številka dovoljenja za promet z zdravilom:
FR/V/8349895 2/1999

Datum spremembe statusa dovoljenja za promet:
13/12/2009

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

Številka postopka:
FR/V/0123/001

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 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 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 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.