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HatchPak Avinew

Ima dovoljenje za promet

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

Informacije o zdravilu

Ime zdravila:
HatchPak Avinew

Učinkovina:
Na voljo samo v English

Ciljne živalske vrste:
Na voljo samo v Spanish Danish Estonian English French Italian Latvian Lithuanian Romanian Finnish Swedish Icelandic Norwegian

Pot uporabe:
Na voljo samo v Spanish Greek English Italian Portuguese

Podatki o zdravilu

Učinkovina / Jakost:
Na voljo samo v English
5.50 log 10 50% embryo infective dose / 1.00 Dose

Farmacevtska oblika:
Inhalacijska suspenzija za nebulator

Karenca glede na pot uporabe:

Coarse spray:

- **Chicken (one day-old chick)**
  - Meat and offal. 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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Na voljo v:

France

Opis ovojnine zdravila:

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 Italian Latvian Norwegian

Imetnik dovoljenja za promet z zdravilom:

Boehringer Ingelheim Animal Health France

Datum dovoljenja za promet z zdravilom:
9/06/2008

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/0792859 2/2008

Datum spremembe statusa dovoljenja za promet:
9/06/2013

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

Številka postopka:
HU/V/105/001

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 Greek 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

Povzetek glavnih značilnosti zdravila

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

Package Leaflet and Labelling

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