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HATCHPAK IB H120 FROZEN SUSPENSION FOR OCULONASAL USE

Ima dovoljenje za promet

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

Informacije o zdravilu

Ime zdravila:
HATCHPAK IB H120 FROZEN SUSPENSION FOR OCULONASAL USE

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:
Okulonazalno dajanje

Podatki o zdravilu

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

Farmacevtska oblika:
Koncentrat za inhalacijsko raztopino za nebulator

Karenca glede na pot uporabe:
Okulonazalno dajanje:

- **Chicken (one day-old chick)**
  - All relevant tissues. 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 Dutch Portuguese Slovak Swedish Icelandic Norwegian

Na voljo v:
Greece

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

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

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

Datum dovoljenja za promet z zdravilom:
15/01/2008

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

Pristojni organ:
National Organization For Medicines

Številka dovoljenja za promet z zdravilom:
16238/27-2-2013/K-0166301

Datum spremembe statusa dovoljenja za promet:
2/06/2021

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

Številka postopka:
FR/V/0171/001

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

Povzetek glavnih značilnosti zdravila

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

Navodilo za uporabo

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

Combined File of all Documents

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