File downloaded on 2026-11-16
Source URL: https://medicines.health.europa.eu/veterinary/sl/600000061502
Nobilis Paramyxo P201 Emulsion for injection

- Pigeon paramyxovirus 1, strain P201, Inactivated

Ima dovoljenje za promet

Informacije o zdravilu

Ime zdravila:
Nobilis Paramyxo P201 Emulsion for injection
U?inkovina:

- Na voljo samo v English

Ciljne živalske vrste:

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

Pot uporabe:

- Subkutana uporaba

Podatki o zdravilu

U?inkovina / Jakost:

- Na voljo samo v English
  6.80
  log2 haemagglutination inhibiting unit(s)
  /
  0.25
  millilitre(s)

Farmacevtska oblika:

- Emulzija za injiciranje

Karenca glede na pot uporabe:

- Subkutana uporaba
  - Pigeon
    - Meat and offal
      0
      day
    - Egg
      0

day

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

- QI01EA01

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

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:

- Intervet Deutschland GmbH

Datum dovoljenja za promet z zdravilom:

- 28/03/2000

Proizvodna mesta, odgovorna za sproš?anje serij:

- Intervet International B.V.

Pristojni organ:

- Paul-Ehrlich-Institut

Številka dovoljenja za promet z zdravilom:

- 243a/97

Datum spremembe statusa dovoljenja za promet:

- 18/01/2010

Referen?na država ?lanica:

- Na voljo samo v Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian

Številka postopka:

- DE/V/0219/001

Zadevne države ?lanice:

- 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
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- Na voljo samo v Estonian English French Lithuanian Portuguese Swedish Icelandic Norwegian

Za poro?ila o domnevnih neželenih u?inkih zdravil za uporabo v veterinaski medicini obiš?ite stran: www.adrreports.eu/vet

Dokumenti

Informacije o zdravilu

Povzetek glavnih zna?ilnosti zdravila

Ta dokument ne obstaja v tem jeziku (slovenš?ina). Lahko ga najdete spodaj v drugem jeziku.
Drugi jeziki (2)
German (PDF)
Objavljeno na dan: 10/02/2022
Updated on: 14/03/2026
Prenesi
English (PDF)
Objavljeno na dan: 28/01/2022
Updated on: 14/03/2026
Prenesi