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Source URL: https://medicines.health.europa.eu/veterinary/en/600000002502

Innovax-ND-IBD (--) - Suspension and solvent for suspension for injection

Ima dovoljenje za promet

- Turkey herpesvirus, strain HVP360 (cell-associated), encoding fusion protein gene of Newcastle disease virus and VP2 protein gene of Infectious bursal disease virus, Live

Informacije o zdravilu

Ime zdravila:
Innovax-ND-IBD (--) - Suspension and solvent for suspension for injection

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 Estonian English French Italian Latvian Lithuanian Hungarian Dutch Romanian Finnish Swedish Icelandic Norwegian

Pot uporabe:
V jajčno celico
Subkutana uporaba

Podatki o zdravilu

Učinkovina / Jakost:
Na voljo samo v English
Presentation_strength:$10^{3,3}$-$10^{4,6}$ PFU Reference:Hse Index:0

Farmacevtska oblika:
Koncentrat in vehikel za suspenzijo za injiciranje

Karenca glede na pot uporabe:
V jajčno celico:

- **Chicken**
  - Not applicable. 0 day Zero days
- **Chicken (embryonated eggs)**
  - Not applicable. 0 day Zero days

Subkutana uporaba:

- **Chicken**
  - Not applicable. 0 day Zero days
- **Chicken (embryonated eggs)**
  - Not applicable. 0 day Zero days

Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):
QI01AD16

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:

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

Na voljo v:

Austria , Belgium , Bulgaria , Cyprus , Czechia , Denmark , France , Germany , Greece , Hungary , Ireland , Italy , Netherlands , Poland , Portugal , Romania , Spain , United Kingdom (Northern Ireland)

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:

Intervet International B.V.

Datum dovoljenja za promet z zdravilom:

22/08/2017

Proizvodna mesta, odgovorna za sproščanje serij:

Intervet International B.V.

Pristojni organ:

European Commission

Številka dovoljenja za promet z zdravilom:

Ta podatek za to zdravilo ni na voljo.

Datum spremembe statusa dovoljenja za promet:
22/08/2017

Za poročila o domnevnih neželenih učinkih zdravil za uporabo v veterinaski medicini obiščite stran: www.adrreports.eu/vet

Dokumenti

Combined File of all Documents

slovenščina (PDF)
Objavljeno na dan: 14/03/2026
Prenesi

ema-puar-v4422-innovax-nd-ibd-ii-0004-en.pdf

ema-puar-innovax-nd-ibd-v-4422-par-en.pdf