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

Atļautas

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

Zāļu identifikācija

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

Aktīvā viela:
Pieejamas tikai English

Mērķsugas:
Cālis
Cāļi (embrionētas olas)

Lietošanas veids:
ievadīšanai olā
subkutānai lietošanai

Sīkāka informācija par zālēm

Aktīvā viela un stiprums:

Pieejamas tikai English
Presentation_strength:$10^{3,3}$-$10^{4,6}$ PFU Reference:Hse Index:0

Farmaceitiskā forma:
Koncentrāts un šķīdinātājs injekciju suspensijas pagatavošanai

Ierobežojumu periods pēc lietošanas veida:
ievadīšanai olā:

- **Cālis**
  - Nav piemērojams. 0 diena Zero days
- **Cāļi (embrionētas olas)**
  - Nav piemērojams. 0 diena Zero days

subkutānai lietošanai:

- **Cālis**
  - Nav piemērojams. 0 diena Zero days
- **Cāļi (embrionētas olas)**
  - Nav piemērojams. 0 diena Zero days

Anatomiski terapeitiski ķīmiskais veterinārais (ATĶvet) kods:
QI01AD16

Izplatīšanas juridiskie nosacījumi:
Recepšu veterinārās zāles

Atļaujas statuss:
Derīga

Atļautas:
Pieejamas tikai Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian

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Pieejamas tikai Estonian English French Lithuanian Portuguese Swedish Icelandic Norwegian

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

Iepakojuma apraksts:
Pieejamas tikai English
Pieejamas tikai English

Papildu informācija

Tiesību tips:

Veterināro zāļu atļaujas piešķiršanas juridiskais pamats:
Pilns pieteikums - jaunai aktīvai vielai (Direktīvas 2001/82/EC 12.(3) pants)

Tirdzniecības atļaujas turētājs:
Intervet International B.V.

Tirdzniecības atļaujas datums:
22/08/2017

Ražošanas vietas sēriju izlaidei:
Intervet International B.V.

Atbildīgā iestāde:
European Commission

Atļaujas numurs:
Šī informācija par šīm zālēm nav pieejama.

Atļaujas statusa maiņas datums:
22/08/2017

Lai uzzinātu par veterināro zāļu blakusparādībām, lūdzu, dodieties uz: www.adrreports.eu/vet

Dokumenti

Visu dokumentu apvienotais fails

latviešu (PDF)
Publicēts: 14/03/2026
Lejupielādēt

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

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