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Ultifend ND IBD (--) - Concentrate and solvent for suspension for injection

Atļautas

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

Zāļu identifikācija

Zāļu nosaukums:
Ultifend ND IBD (--) - Concentrate 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:4000 PFU/dose - 12000 PFU/dose Index:0

Farmaceitiskā forma:
Koncentrāts un šķīdinātājs injekciju suspensijas pagatavošanai

Ierobežojumu periods pēc lietošanas veida:
ievadīšanai olā:

- **Cālis**
  - Nav piemērojams. 0 diena Zero days

subkutānai lietošanai:

- **Cālis**
  - Nav piemērojams. 0 diena Zero days

Anatomiski terapeitiski ķīmiskais veterinārais (ATĶvet) kods:
QI01AD16

Izplatīšanas juridiskie nosacījumi:
Recepšu veterinārās zāles

Atļaujas statuss:
Derīga

Atļautas:
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Iepakojuma apraksts:

Pieejamas tikai English

Pieejamas tikai English

Pieejamas tikai English

Papildu informācija

Tiesību tips:

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

Tirdzniecības atļaujas turētājs:
Ceva-Phylaxia Zrt.

Tirdzniecības atļaujas datums:
20/04/2021

Ražošanas vietas sēriju izlaidei:
Ceva-Phylaxia Zrt.

Atbildīgā iestāde:
European Commission

Atļaujas numurs:
Šī informācija par šīm zālēm nav pieejama.

Atļaujas statusa maiņas datums:
20/04/2021

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