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Nobivac NXT HCPFeLV (--) + (--) + (--) + (--) - Lyophilisate and solvent for suspension for injection

Atļautas

- Felid herpesvirus 1, strain G2620A, Live
- Feline calicivirus, strain F9, Live
- Feline panleucopenia virus, strain MW-1, Live
- Feline leukemia virus (strain A/Glasgow-1), gp85 gene encoded by Venezuelan equine encephalitis virus (strain TC-83), self-amplifying RNA, viral particle, replication deficient

Zāļu identifikācija

Zāļu nosaukums:
Nobivac NXT HCPFeLV (--) + (--) + (--) + (--) - Lyophilisate and solvent for suspension for injection

Aktīvā viela:
Pieejamas tikai English
Pieejamas tikai English
Pieejamas tikai English
Pieejamas tikai English

Mērķsugas:
Kaķis

Lietošanas veids:
subkutānai lietošanai

Sīkāka informācija par zālēm

Aktīvā viela un stiprums:
Pieejamas tikai English
Presentation_strength:$10^{5,8}$ – $10^{7,1}$ PFU Index:0
Pieejamas tikai English
Presentation_strength:$10^{4,6}$ – $10^{7,0}$ PFU Index:1
Pieejamas tikai English
Presentation_strength:$10^{5,2}$ – $10^{6,4}$ $TCID_{50}$ Index:2
Pieejamas tikai English
Presentation_strength:$10^{6,6}$ – $10^{9,3}$ RP Index:3

Farmaceitiskā forma:
Liofilizāts un šķīdinātājs suspensijas injekcijām pagatavošanai

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Anatomiski terapeitiski ķīmiskais veterinārais (ATĶvet) kods:
QI06AX

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

Iepakojuma apraksts:

Pieejamas tikai English

Pieejamas tikai English

Papildu informācija

Tiesību tips:

Veterināro zāļu atļaujas piešķiršanas juridiskais pamats:

Pilns pieteikums - jaunai aktīvai vielai (ES Regulas 2019/6 8.pants)

Tirdzniecības atļaujas turētājs:
Intervet International B.V.

Tirdzniecības atļaujas datums:
29/07/2026

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:
29/07/2026

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

Dokumenti

Visu dokumentu apvienotais fails

latviešu (PDF)
Publicēts: 25/08/2026
Lejupielādēt

ema-puar-v6682-nobivacnxthcpfelv-initial-en.pdf