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

Ima dovoljenje za promet

- 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

Informacije o zdravilu

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

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Učinkovina:
Na voljo samo v English
Na voljo samo v English
Na voljo samo v English
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

Pot uporabe:
Subkutana uporaba

Podatki o zdravilu

Učinkovina / Jakost:
Na voljo samo v English
Presentation_strength:$10^{5,8}$ – $10^{7,1}$ PFU Index:0
Na voljo samo v English
Presentation_strength:$10^{4,6}$ – $10^{7,0}$ PFU Index:1
Na voljo samo v English
Presentation_strength:$10^{5,2}$ – $10^{6,4}$ $TCID_{50}$ Index:2
Na voljo samo v English
Presentation_strength:$10^{6,6}$ – $10^{9,3}$ RP Index:3

Farmacevtska oblika:
Liofilizat in vehikel za suspenzijo za injiciranje

Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):
QI06AX

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

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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 French Italian Latvian Lithuanian Norwegian

Imetnik dovoljenja za promet z zdravilom:
Intervet International B.V.

Datum dovoljenja za promet z zdravilom:
29/07/2026

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:
29/07/2026

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: 25/08/2026
Prenesi

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