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Nobivac Myxo-RHD Plus (--) - Lyophilisate and solvent for suspension for injection

Ima dovoljenje za promet

- Myxoma virus, strain 009, expressing capsid protein gene of Rabbit haemorrhagic disease virus, Live
- Myxoma virus, strain MK1899, expressing capsid protein gene of Rabbit haemorrhagic disease virus 2, Live

Informacije o zdravilu

Ime zdravila:
Nobivac Myxo-RHD Plus (--) - Lyophilisate and solvent for suspension for injection

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

Ciljne živalske vrste:
Na voljo samo v Bulgarian Spanish Czech Danish 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:≥$10^{3,0}$ and ≤$10^{5,8}$ FFU Comments:2.c.2.1 Index:0
Na voljo samo v English
Presentation_strength:≥$10^{3,0}$ and ≤$10^{5,8}$ FFU Comments:2.c.2.1 Index:1

Farmacevtska oblika:
Liofilizat in vehikel za suspenzijo za injiciranje

Karenca glede na pot uporabe:
Subkutana uporaba:

- **Rabbit**
  - Not applicable. 0 day Zero days

Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):
QI08AD

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:
Na voljo samo v Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian
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Na voljo samo v Estonian English French Lithuanian Portuguese Swedish Icelandic Norwegian

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Na voljo v:
Austria , Belgium , Czechia , Denmark , Finland , France , Germany , Greece , Hungary , Ireland , Italy , Luxembourg , Netherlands , Poland , Portugal , Slovakia , Spain , Sweden , United Kingdom (Northern Ireland)

Opis ovojnine zdravila:

Na voljo samo v English
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:
19/11/2019

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:
19/11/2019

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-nobivac-myxo-rhd-plus-v-4989-par-en.pdf