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RP Vacc emulsion for injection for pigeons

Ima dovoljenje za promet

- Pigeon paramyxovirus 1, strain 988M, Inactivated
- Pigeon rotavirus A, strain Ro/D, Inactivated

Informacije o zdravilu

Ime zdravila:
RP Vacc emulsion for injection for pigeons

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

Ciljne živalske vrste:
Na voljo samo v Bulgarian Spanish Czech Danish German Estonian English French Italian Latvian Lithuanian Hungarian Dutch Romanian Finnish Swedish Icelandic Norwegian

Pot uporabe:
intramuskularna uporaba
Subkutana uporaba

Podatki o zdravilu

Učinkovina / Jakost:

Na voljo samo v English
6.47 log2 haemagglutination inhibiting unit(s) / 0.30 millilitre(s)

Na voljo samo v English
52.20 enzyme-linked immunosorbent assay unit / 0.30 millilitre(s)

Farmacevtska oblika:

Emulzija za injiciranje

Karenca glede na pot uporabe:

intramuskularna uporaba:

- **Pigeon**
  - All relevant tissues. 0 day — Zero days

Subkutana uporaba:

- **Pigeon**
  - All relevant tissues. 0 day

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

QI01EA

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

Opis ovojnine zdravila:

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

Imetnik dovoljenja za promet z zdravilom:
Pharmagal Bio spol. s r.o.

Datum dovoljenja za promet z zdravilom:
6/12/2021

Proizvodna mesta, odgovorna za sproščanje serij:
Pharmagal Bio spol. s r.o.

Pristojni organ:
Directorate Of Veterinary Medicinal Products

Številka dovoljenja za promet z zdravilom:
4293/X/21 NÉBIH ÁTI

Datum spremembe statusa dovoljenja za promet:
6/12/2021

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

Številka postopka:
SK/V/0110/001

Zadevne države članice:

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

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Za poročila o domnevnih neželenih učinkih zdravil za uporabo v veterinaski medicini obiščite stran: www.adrreports.eu/vet

Dokumenti

Combined File of all Documents

Ta dokument ne obstaja v tem jeziku (slovenščina). Lahko ga najdete spodaj v drugem jeziku.