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RABADROP, peroralna suspenzija

Ima dovoljenje za promet

- Rabies virus, strain SAD, Live

Informacije o zdravilu

Ime zdravila:
RABADROP, peroralna suspenzija

Učinkovina:
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 Norwegian rakunasti pes

Pot uporabe:
Peroralna uporaba

Podatki o zdravilu

Učinkovina / Jakost:
Na voljo samo v English
8.50 log10 50% tissue culture infectious dose / 1.00 Dose

Farmacevtska oblika:

Peroralna suspenzija

Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):
QI07BD

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
Na voljo samo v English
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

Imetnik dovoljenja za promet z zdravilom:
Bioveta a.s.

Datum dovoljenja za promet z zdravilom:
1/07/2019

Proizvodna mesta, odgovorna za sproščanje serij:

Bioveta a.s.

Pristojni organ:
Agency For Medicinal Products And Medical Devices Of The Republic Of Slovenia

Številka dovoljenja za promet z zdravilom:
DC/V/0672/001

Datum spremembe statusa dovoljenja za promet:
1/07/2019

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

Številka postopka:
CZ/V/0149/001

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 Croatian Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian
Na voljo samo v Spanish German Estonian English French Italian Dutch Portuguese Slovak Finnish 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

Povzetek glavnih značilnosti zdravila

slovenščina (PDF)
Objavljeno na dan: 3/03/2022
Updated on: 14/03/2026
Prenesi

Navodilo za uporabo

slovenščina (PDF)
Objavljeno na dan: 3/03/2022
Updated on: 14/03/2026
Prenesi

Označevanje

slovenščina (PDF)
Objavljeno na dan: 3/03/2022
Updated on: 14/03/2026
Prenesi

eu-puar-czv0149001-mr-rabadrop-en.pdf