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Cerenia 10 mg/ml - Solution for injection

Ima dovoljenje za promet

- Maropitant citrate

Informacije o zdravilu

Ime zdravila:
Cerenia 10 mg/ml - Solution for injection

Učinkovina:
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
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:
Intravenska uporaba
Subkutana uporaba

Podatki o zdravilu

Učinkovina / Jakost:

Na voljo samo v English
14.49 milligram(s) / 1.00 Viala

Farmacevtska oblika:

Raztopina za injiciranje

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

QA04AD90

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 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

Na voljo v:
Austria , Belgium , Bulgaria , Czechia , Denmark , Estonia , Finland , Germany , Hungary , Ireland , Italy , Latvia , Lithuania , Luxembourg , Netherlands , Poland , Romania , Slovakia , Spain , Sweden , United Kingdom (Northern Ireland)

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

Imetnik dovoljenja za promet z zdravilom:
Zoetis Belgium

Datum dovoljenja za promet z zdravilom:
29/09/2006

Proizvodna mesta, odgovorna za sproščanje serij:
Zoetis Manufacturing & Research Spain S.L.

Pristojni organ:
European Commission

Številka dovoljenja za promet z zdravilom:
Ta podatek za to zdravilo ni na voljo.

Datum spremembe statusa dovoljenja za promet:
29/09/2006

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-cerenia-v-106-var-x-0023-en.pdf

ema-puar-cerenia-v-106-par-en.pdf

ema-puar-cerenia-v-106-var-ii-0013-en.pdf