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COFACALCIUM, injekcinis tirpalas

Ni odobreno

- Magnesium hypophosphite hexahydrate
- Calcium gluconate

Informacije o zdravilu

Ime zdravila:
COFACALCIUM, injekcinis tirpalas

Učinkovina:
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
govedo
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
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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
intramuskularna uporaba
Subkutana uporaba

Podatki o zdravilu

Učinkovina / Jakost:
Na voljo samo v English
90.00 milligram(s) / 1.00 millilitre(s)

Na voljo samo v English
280.00 milligram(s) / 1.00 millilitre(s)

Farmacevtska oblika:
Raztopina za injiciranje

Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):
QA12AX

Pravni status za dobavo/izdajo zdravila:
Na voljo samo v English French Italian Latvian Lithuanian Portuguese Finnish Swedish Norwegian

Status dovoljenja za promet z zdravilom:
Surrendered

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 Lithuanian

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 Norwegian

Imetnik dovoljenja za promet z zdravilom:
Dopharma Research B.V.

Datum dovoljenja za promet z zdravilom:
21/09/2010

Proizvodna mesta, odgovorna za sproščanje serij:
Dopharma France

Pristojni organ:
State Food And Veterinary Service

Številka dovoljenja za promet z zdravilom:
LT/2/10/1971/002

Datum spremembe statusa dovoljenja za promet:
6/10/2015

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

Dokumenti

RV1971.pdf