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C-B-GLUCONAT 38 % PLUS 6 % infuzinis tirpalas arkliams, galvijams, avims, ožkoms ir kiaulėms

Ni odobreno

- Boric acid
- Magnesium chloride hexahydrate
- Calcium gluconate

Informacije o zdravilu

Ime zdravila:
C-B-GLUCONAT 38 % PLUS 6 % infuzinis tirpalas arkliams, galvijams, avims, ožkoms ir kiaulėms

Učinkovina:
Na voljo samo v English
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
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

Podatki o zdravilu

Učinkovina / Jakost:
Na voljo samo v English
5.00 gram(s) / 100.00 millilitre(s)

Na voljo samo v English
6.00 gram(s) / 100.00 millilitre(s)

Na voljo samo v English
38.00 gram(s) / 100.00 millilitre(s)

Farmacevtska oblika:
Raztopina za infundiranje

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
Na voljo samo v Lithuanian
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 Italian Latvian Lithuanian Norwegian

Imetnik dovoljenja za promet z zdravilom:
Bela-Pharm GmbH & Co. KG

Datum dovoljenja za promet z zdravilom:
25/03/2007

Proizvodna mesta, odgovorna za sproščanje serij:
Bela-Pharm GmbH & Co. KG

Pristojni organ:
State Food And Veterinary Service

Številka dovoljenja za promet z zdravilom:
LT/2/07/1736/001

Datum spremembe statusa dovoljenja za promet:
29/03/2012

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

Dokumenti

RV1736.pdf