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Calcitat N 25 pro inf.

Ima dovoljenje za promet

- Calcium gluconate monohydrate
- Magnesium chloride hexahydrate
- Phosphorylcolamine
- Calcium hydroxide
- Calcium borogluconate

Informacije o zdravilu

Ime zdravila:
Calcitat N 25 pro inf.

Učinkovina:
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

Ciljne živalske vrste:
govedo
tele
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
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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

Podatki o zdravilu

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

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

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

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

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

Farmacevtska oblika:
Raztopina za infundiranje

Karenca glede na pot uporabe:
Intravenska uporaba:

-

govedo
- Meat and offal. 0 day

•
tele
- Meat and offal. 0 day

•
Goat
- Meat and offal. 0 day

•
Sheep
- Meat and offal. 0 day

•
Pig
- Meat and offal. 0 day

•
Pig (piglet)
- Meat and offal. 0 day

Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):
QA12AA20

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 v:
Germany

Opis ovojnine zdravila:
Na voljo samo v German
Na voljo samo v German
Na voljo samo v German
Na voljo samo v German
Na voljo samo v German
Na voljo samo v German

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:
aniMedica GmbH

Datum dovoljenja za promet z zdravilom:
30/03/1981

Proizvodna mesta, odgovorna za sproščanje serij:
Pasteur Filiala Filipesti S.A.

Pristojni organ:
Federal Office Of Consumer Protection And Food Safety

Številka dovoljenja za promet z zdravilom:
1468.00.00

Datum spremembe statusa dovoljenja za promet:
9/11/2009

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.