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LACHESIS COMPOSITUM N, injekcinis tirpalas

Ima dovoljenje za promet

- LACHESIS MUTUS DIL D6
- PYROGENIUM NOSODE D6
- JUNIPERUS SABINA D3
- ECHINACEA ANGUSTIFOLIA
- PULSATILLA PRATENSIS D2

Informacije o zdravilu

Ime zdravila:
LACHESIS COMPOSITUM N, injekcinis tirpalas

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:
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

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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
psice
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:
Subkutana uporaba

Podatki o zdravilu

Učinkovina / Jakost:
Na voljo samo v English
200.00 milligram(s) / 1.00 Ampula

Na voljo samo v English
200.00 milligram(s) / 1.00 Ampula

Na voljo samo v English
100.00 milligram(s) / 1.00 Ampula

Na voljo samo v English
100.00 milligram(s) / 1.00 Ampula

Na voljo samo v English
50.00 milligram(s) / 1.00 Ampula

Farmacevtska oblika:
Raztopina za injiciranje

Pravni status za dobavo/izdajo zdravila:
Ta podatek za to zdravilo ni na voljo.

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 Lithuanian

Na voljo samo v Lithuanian

Dodatne informacije

Vrsta dovoljenja:

Na voljo samo v English French Italian Latvian Swedish Icelandic Norwegian

Imetnik dovoljenja za promet z zdravilom:

Biologische Heilmittel Heel GmbH

Datum dovoljenja za promet z zdravilom:

23/03/2003

Proizvodna mesta, odgovorna za sproščanje serij:

Biologische Heilmittel Heel GmbH

Pristojni organ:

State Food And Veterinary Service

Številka dovoljenja za promet z zdravilom:

LT/2/03/1535/001-002

Datum spremembe statusa dovoljenja za promet:

25/03/2008

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

Dokumenti

Navodilo za uporabo

Ta dokument ne obstaja v tem jeziku (slovenščina). Lahko ga najdete spodaj v drugem jeziku.