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Echinacea compositum ad us. vet.

Ima dovoljenje za promet

- HYDRARGYRUM BICHLORATUM D6
- PHOSPHORUS D8
- ACONITUM NAPELLUS D4
- BRYONIA D6
- ARNICA MONTANA D6
- LACHESIS D10
- ECHINACEA D3
- SULFUR D8

Informacije o zdravilu

Ime zdravila:
Echinacea compositum ad us. vet.

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
Na voljo samo v English
Na voljo samo v English
Na voljo samo v English

Ciljne živalske vrste:
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
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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:
Subkutana uporaba

Podatki o zdravilu

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

Na voljo samo v English
0.10 gram(s) / 5.00 millilitre(s)

Na voljo samo v English
0.10 gram(s) / 5.00 millilitre(s)

Na voljo samo v English
0.10 gram(s) / 5.00 millilitre(s)

Na voljo samo v English
0.10 gram(s) / 5.00 millilitre(s)

Na voljo samo v English
0.10 gram(s) / 5.00 millilitre(s)

Na voljo samo v English
0.50 gram(s) / 5.00 millilitre(s)

Na voljo samo v English
0.10 gram(s) / 5.00 millilitre(s)

Farmacevtska oblika:

Raztopina za injiciranje

Pravni status za dobavo/izdajo zdravila:

Zdravilo, ki se izdaja brez veterinarskega recepta

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 German

Na voljo samo v German

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:

28/01/1993

Proizvodna mesta, odgovorna za sproščanje serij:

Biologische Heilmittel Heel GmbH

Pristojni organ:

Federal Office Of Consumer Protection And Food Safety

Številka dovoljenja za promet z zdravilom:

17758.00.00

Datum spremembe statusa dovoljenja za promet:
25/03/2010

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.