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Phytolacca S-logoplex Flüssige Verdünnung zur Injektion für Pferde, Rinder, Kälber, Schweine, Ferkel, Schafe, Ziegen, Hunde, Katzen, Kaninchen, Kleinnager und Frettchen

Ima dovoljenje za promet

- Phytolacca americana D5
- ACIDUM FORMICICUM D4
- ACIDUM SILICICUM D10
- ARNICA MONTANA D6
- ATROPA BELLA-DONNA D6
- CALCIUM FLUORATUM D8
- CALCIUM SULFURICUM D12
- HEPAR SULFURIS D8
- LOPHOPHYTUM LEANDRI DIL. D6

Informacije o zdravilu

Ime zdravila:
Phytolacca S-logoplex Flüssige Verdünnung zur Injektion für Pferde, Rinder, Kälber, Schweine, Ferkel, Schafe, Ziegen, Hunde, Katzen, Kaninchen, Kleinnager und Frettchen

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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
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

Pot uporabe:
Subkutana uporaba

Podatki o zdravilu

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

Na voljo samo v English
10.00 gram(s) / 100.00 gram(s)

Na voljo samo v English
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10.00 gram(s) / 100.00 gram(s)

Na voljo samo v English
10.00 gram(s) / 100.00 gram(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:
ZIEGLER GmbH Homoeopathika ad us. vet.

Datum dovoljenja za promet z zdravilom:
1/04/2023

Proizvodna mesta, odgovorna za sproščanje serij:
ZIEGLER GmbH Homoeopathika ad us. vet.

Pristojni organ:
Federal Office Of Consumer Protection And Food Safety

Številka dovoljenja za promet z zdravilom:
V7008585.00.00

Datum spremembe statusa dovoljenja za promet:
1/04/2023

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.