File downloaded on 2026-11-05
Source URL: https://medicines.health.europa.eu/veterinary/sl/600000100032

Flor de piedra-logoplex

Ima dovoljenje za promet

- STRYCHNOS NUX-VOMICA D6
- LOPHOPHYTUM LEANDRI D3
- LACHESIS MUTUS DIL. D8
- CHELIDONIUM MAJUS D4

Informacije o zdravilu

Ime zdravila:
Flor de piedra-logoplex

Učinkovina:
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

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:
Subkutana uporaba

Podatki o zdravilu

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

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

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

Na voljo samo v English
0.25 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:
ZIEGLER VetProducts GmbH

Datum dovoljenja za promet z zdravilom:
12/05/2006

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:
131430.00.00

Datum spremembe statusa dovoljenja za promet:
19/07/2013

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.