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Ni odobreno

- CYPRIPEDIUM PARVIFLORUM VAR. PUBESCENS D8
- ORIGANUM MAJORANA D4
- AQUILEGIA VULGARIS D4
- MOSCHUS MOSCHIFERUS D6
- CYCLAMEN PURPURASCENS D4
- CONYZA CANADENSIS D3
- SENECIO NEMORENSIS D3
- CAPSELLA BURSA-PASTORIS D3
- MYRISTICA FRAGRANS D6
- STRYCHNOS IGNATII D6
- CALCIUM CARBONICUM HAHNEMANNI D8
- SEPIA OFFICINALIS D6
- VIBURNUM OPULUS D3
- ACIDUM NITRICUM D4
- PULSATILLA PRATENSIS D4

Informacije o zdravilu

Ime zdravila:
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Učinkovina:
Na voljo samo v English
Na voljo samo v English
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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

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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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Pot uporabe:

Subkutana uporaba

Podatki o zdravilu

Učinkovina / Jakost:

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

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

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

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

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

Na voljo samo v English
0.00 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:
Surrendered

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:
24/07/1993

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:
23271.00.00

Datum spremembe statusa dovoljenja za promet:
5/08/2026

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.