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Cactus compositum ad us. vet.

Ima dovoljenje za promet

- SELENICEREUS GRANDIFLORUS D3
- CRATAEGUS D2
- SPIGELIA ANTHELMIA D5
- KALIUM CARBONICUM D5
- NITROGLYCERINUM D5

Informacije o zdravilu

Ime zdravila:
Cactus 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

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
50.00 milligram(s) / 5.00 millilitre(s)

Na voljo samo v English
50.00 milligram(s) / 5.00 millilitre(s)

Na voljo samo v English
50.00 milligram(s) / 5.00 millilitre(s)

Na voljo samo v English
50.00 milligram(s) / 5.00 millilitre(s)

Na voljo samo v English
30.00 milligram(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:
31/10/1983

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:
4186.00.00

Datum spremembe statusa dovoljenja za promet:
15/08/2013

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.