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Dr. Reckeweg ReVet RV 5

Ima dovoljenje za promet

- Lophophytum leandri C2
- Okoubaka aubrevillei C2
- Silybum marianum C2

Informacije o zdravilu

Ime zdravila:
Dr. Reckeweg ReVet RV 5

Učinkovina:
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
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
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
govedo

Pot uporabe:
Peroralna uporaba

Podatki o zdravilu

Učinkovina / Jakost:
Na voljo samo v English
3.33 milligram(s) / 1.00 gram(s)

Na voljo samo v English
3.33 milligram(s) / 1.00 gram(s)

Na voljo samo v English
3.33 milligram(s) / 1.00 gram(s)

Farmacevtska oblika:
Zrnca

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 Swedish

Dodatne informacije

Vrsta dovoljenja:
Na voljo samo v English French Italian Latvian Swedish Icelandic Norwegian

Imetnik dovoljenja za promet z zdravilom:
Biosan AB

Datum dovoljenja za promet z zdravilom:
16/12/2004

Proizvodna mesta, odgovorna za sproščanje serij:
Pharmazeutische Fabrik Dr. Reckeweg & Co. GmbH

Pristojni organ:
Swedish Medical Products Agency

Številka dovoljenja za promet z zdravilom:
37998

Datum spremembe statusa dovoljenja za promet:
16/12/2004

Za poročila o domnevnih neželenih učinkih zdravil za uporabo v veterinaski medicini obiščite stran: www.adrreports.eu/vet