

ReVet RV 18

Pooblašeno

- Berberis vulgaris C4
- LYTTA VESICATORIA C6
- LYCOPODIUM CLAVATUM C9
- Sabal serrulata C4

Product identification

Ime zdravila:

ReVet RV 18

Učinkovina:

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](#) [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

Na voljo samo v [Bulgarian](#) [Spanish](#) [Czech](#) [Danish](#) [Estonian](#) [Greek](#) [English](#) [French](#) [Italian](#) [Latvian](#) [Lithuanian](#) [Hungarian](#) [Dutch](#) [Romanian](#) [Finnish](#) [Swedish](#) [Norwegian](#) [plazilec](#)

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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Norwegian

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Na voljo samo v Bulgarian Spanish Danish German Estonian English French Italian Latvian Lithuanian Hungarian Dutch Romanian Finnish Swedish Norwegian

Pot uporabe:

Peroralna uporaba

Product details

Učinkovina / Jakost:

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

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

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

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

Farmacevtska oblika:

Kroglice

Pravni status za dobavo / izdajo zdravila:

Zdravilo, ki se izdaja brez veterinarskega recepta

Status dovoljenja:

Valid

Authorised in:

Na voljo samo v Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian

Opis ovojnine:

Na voljo samo v German

Na voljo samo v [German](#)
Na voljo samo v [German](#)
Na voljo samo v [German](#)

Additional information

Entitlement type:

Na voljo samo v [English](#) [French](#) [Italian](#) [Latvian](#) [Swedish](#) [Icelandic](#) [Norwegian](#)

Imetnik dovoljenja za promet z zdravilom:

Pharmazeutische Fabrik Dr. Reckeweg & Co. GmbH

Marketing authorisation date:

10/03/2017

Proizvodna mesta, odgovorna za sproščanje serij:

Pharmazeutische Fabrik Dr. Reckeweg & Co. GmbH

Pristojni organ:

Federal Office Of Consumer Protection And Food Safety

Številka dovoljenja :

402335.00.00

Datum spremembe statusa dovoljenja:

1/11/2021

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

Documents

Povzetek glavnih značilnosti zdravila

Ta dokument ne obstaja v tem jeziku (slovenščina). Spodaj ga najdete v drugem jeziku.

Navodilo za uporabo

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