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NeyDIL 65 Revitorgan-Dilution Nr.65 D4 pro vet.

Ni odobreno

- Thymus bovis fetalis D4 aquos.
- Glandula suprarenalis suis D4 aquos.
- Nodus lymphaticus bovis fetalis D4 aquos.
- Glandula parathyreoidea suis D4 aquos.
- Hepar bovis fetalis D4 aquos.
- Hepar bovis D4 aquos.
- Ren bovis fetalis D4 aquos.
- Ren bovis fetalis D4 aquos.
- Pancreas bovis D4 aquos.
- Lien bovis fetalis D4 aquos.
- Vasa bovis fetalis D4 aquos.
- Funiculus umbilicalis bovis fetalis D4 aquos.
- Hypophysis cerebri suis D4 aquos.
- Diencephalon bovis fetalis D4 aquos.
- Mucosa miscae bovis D4 aquos.
- Cutis bovis fetalis D4 aquos.
- Medulla ossium bovis fetalis D4 aquos.
- Mucosa miscae bovis fetalis D4 aquos.

Informacije o zdravilu

Ime zdravila:
NeyDIL 65 Revitorgan-Dilution Nr.65 D4 pro vet.

Učinkovina:

Na voljo samo v English
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Ciljne živalske vrste:

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

Na voljo samo v Bulgarian Spanish Czech Danish German Estonian Greek English French Italian Latvian Lithuanian Hungarian Romanian Finnish Swedish 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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Pot uporabe:

intramuskularna uporaba

Subkutana uporaba

Podatki o zdravilu

Učinkovina / Jakost:

Na voljo samo v English
1.00 unit(s) / 2.00 millilitre(s)

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1.00 unit(s) / 2.00 millilitre(s)

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1.00 unit(s) / 2.00 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:
Vitorgan Arzneimittel GmbH

Datum dovoljenja za promet z zdravilom:
22/08/2005

Proizvodna mesta, odgovorna za sproščanje serij:
Vitorgan Arzneimittel GmbH
Solupharm Pharmazeutische Erzeugnisse GmbH

Pristojni organ:
Federal Office Of Consumer Protection And Food Safety

Številka dovoljenja za promet z zdravilom:

2580391.00.00

Datum spremembe statusa dovoljenja za promet:
22/01/2026

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