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BERBERIS HOMACCORD

Ima dovoljenje za promet

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- CITRULLUS COLOCYNTHIS D200
- CITRULLUS COLOCYNTHIS D30
- CITRULLUS COLOCYNTHIS D4
- BERBERIS VULGARIS D10
- BERBERIS VULGARIS D200
- BERBERIS VULGARIS D4
- VERATRUM ALBUM D10
- VERATRUM ALBUM D200
- VERATRUM ALBUM D30
- VERATRUM ALBUM D5

Informacije o zdravilu

Ime zdravila:
BERBERIS HOMACCORD

Učinkovina:
Na voljo samo v English
Na voljo samo v English
Na voljo samo v English
Na voljo samo v English
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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

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Pot uporabe:

Peroralna uporaba

intramuskularna uporaba

Intravenska uporaba

Subkutana uporaba

Podatki o zdravilu

Učinkovina / Jakost:

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

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

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

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

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

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

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

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

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

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

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

Farmacevtska oblika:

Raztopina za injiciranje

Pravni status za dobavo/izdajo zdravila:

Zdravilo, ki se izdaja na veterinarski recept

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 Italian
Na voljo samo v Italian

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:
24/12/2014

Proizvodna mesta, odgovorna za sproščanje serij:
Biologische Heilmittel Heel GmbH

Pristojni organ:
Ministry Of Health

Številka dovoljenja za promet z zdravilom:
Ta podatek za to zdravilo ni na voljo.

Datum spremembe statusa dovoljenja za promet:
24/12/2014

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

Dokumenti

Package Leaflet and Labelling

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