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CHELIDONIUM -HOMACCORD VETERINARIO

Ima dovoljenje za promet

- ATROPA BELLA-DONNA D10
- ATROPA BELLA-DONNA D1000
- ATROPA BELLA-DONNA D200
- ATROPA BELLA-DONNA D30
- ATROPA BELLA-DONNA D4
- CHELIDONIUM MAJUS D10
- CHELIDONIUM MAJUS D200
- FEL TAURI D10
- FEL TAURI D200

Informacije o zdravilu

Ime zdravila:
CHELIDONIUM -HOMACCORD VETERINARIO

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
Na voljo samo v English
Na voljo samo v English
Na voljo samo v English
Na voljo samo v English

Ciljne živalske vrste:

krava

govedo

okrasna ptica

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:

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
15.00 milligram(s) / 5.00 millilitre(s)

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

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

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

Na voljo samo v English
5.00 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:
30/10/2012

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:
4/10/2012

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.