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SOLUCLIN 25 MG/ML ORAL SOLUTION FOR CATS AND DOGS

Ima dovoljenje za promet

- Clindamycin hydrochloride

Informacije o zdravilu

Ime zdravila:
SOLUCLIN 25 MG/ML ORAL SOLUTION FOR CATS AND DOGS

Učinkovina:
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

Pot uporabe:
Peroralna uporaba

Podatki o zdravilu

Učinkovina / Jakost:
Na voljo samo v English
27.15 milligram(s) / 1.00 millilitre(s)

Farmacevtska oblika:
Peroralna raztopina

Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):
QJ01FF01

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

Na voljo v:
Denmark

Opis ovojnine zdravila:
Na voljo samo v English
Na voljo samo v English
Na voljo samo v English

Dodatne informacije

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

Pravna podlaga za izdajo dovoljenja za promet z zdravilom:

Na voljo samo v English French Italian Latvian Lithuanian Norwegian

Imetnik dovoljenja za promet z zdravilom:
CP-Pharma Handelsgesellschaft mbH

Datum dovoljenja za promet z zdravilom:
26/06/2023

Proizvodna mesta, odgovorna za sproščanje serij:
CP-Pharma Handelsgesellschaft mbH

Pristojni organ:
Danish Medicines Agency

Številka dovoljenja za promet z zdravilom:
67613

Datum spremembe statusa dovoljenja za promet:
26/06/2023

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

Številka postopka:
FR/V/0455/001

Zadevne države članice:
Na voljo samo v Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian
Na voljo samo v Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian
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Na voljo samo v Estonian English French Lithuanian Portuguese Swedish Icelandic Norwegian

Generic of:
600000039971

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

Dokumenti

Povzetek glavnih značilnosti zdravila

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

Combined File of all Documents

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