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Source URL: https://medicines.health.europa.eu/veterinary/en/600000000067

Equioxx 57 mg - Chewable tablet

Ima dovoljenje za promet

- Firocoxib

Informacije o zdravilu

Ime zdravila:
Equioxx 57 mg - Chewable tablet

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

Pot uporabe:
Peroralna uporaba

Podatki o zdravilu

Učinkovina / Jakost:
Na voljo samo v English
57.00 milligram(s) / 1.00 Tableta

Farmacevtska oblika:

Žvečljiva tableta

Karenca glede na pot uporabe:

Peroralna uporaba:

- **Horse**
  - Meat and offal. 26 day 26 days

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

QM01AH90

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

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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

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Na voljo v:

Austria , Belgium , Denmark , France , Germany , Ireland , Italy , Liechtenstein , Luxembourg , Malta , Netherlands , Norway , Poland , Spain , Sweden ,

United Kingdom (Northern Ireland)

Opis ovojnine zdravila:

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

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 Italian Latvian Norwegian

Imetnik dovoljenja za promet z zdravilom:
Audevard

Datum dovoljenja za promet z zdravilom:
25/06/2008

Proizvodna mesta, odgovorna za sproščanje serij:
Boehringer Ingelheim Animal Health France
PROVET S.A.

Pristojni organ:
European Commission

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

Datum spremembe statusa dovoljenja za promet:
9/02/2017

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

Dokumenti

Combined File of all Documents

slovenščina (PDF)
Objavljeno na dan: 10/04/2026
Prenesi

ema-puar-equioxx-v-142-var-x-0015-en.pdf

ema-puar-equioxx-v-142-par-en.pdf

RV0838.pdf