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Quinox 60.6 mg/g Premix for Medicated Feeding Stuff for Cattle and Sheep

Ima dovoljenje za promet

- Decoquinate

Informacije o zdravilu

Ime zdravila:
Quinox 60.6 mg/g Premix for Medicated Feeding Stuff for Cattle and Sheep

Učinkovina:
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

Pot uporabe:
Peroralna uporaba

Podatki o zdravilu

Učinkovina / Jakost:
Na voljo samo v English

60.60 milligram(s) / 1.00 gram(s)

Farmacevtska oblika:
Predmešanica za zdravilno krmno mešanico

Karenca glede na pot uporabe:
Peroralna uporaba:

- **govedo**
  - Meat and offal. 0 day
- **Sheep**
  - Meat and offal. 0 day

Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):
QP51BX04

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 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:
HuVepharma

Datum dovoljenja za promet z zdravilom:
2/09/2026

Proizvodna mesta, odgovorna za sproščanje serij:
Huvepharma S.A.

Pristojni organ:
National Organization For Medicines

Številka dovoljenja za promet z zdravilom:
103976/03-09-2026/10-09-2026/K-0265201

Datum spremembe statusa dovoljenja za promet:
30/06/2026

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

Številka postopka:
IE/V/0589/001

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 Greek English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian
Na voljo samo v Spanish Czech German Estonian English French Italian Latvian Dutch Portuguese Slovak Swedish Icelandic Norwegian
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Za poročila o domnevnih neželenih učinkih zdravil za uporabo v veterinaski medicini obiščite stran: www.adrreports.eu/vet