File downloaded on 2026-11-08
**Source URL:** https://medicines.health.europa.eu/veterinary/sl/700000225726

# Quinox 6 mg/g oral powder for cattle and sheep

Ima dovoljenje za promet

- Decoquinate

## Informacije o zdravilu

**Ime zdravila:**
Quinox 6 mg/g oral powder 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
6.00 milligram(s) / 1.00 gram(s)

**Farmacevtska oblika:**
Peroralni prašek

**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:**
28/07/2026

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

**Pristojni organ:**
Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

**Številka dovoljenja za promet z zdravilom:**
3521

**Datum spremembe statusa dovoljenja za promet:**
28/07/2026

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

**Številka postopka:**
IE/V/0599/001

**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
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

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
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
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
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 Croatian Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian
Na voljo samo v Estonian English French Lithuanian Portuguese Swedish Icelandic Norwegian

---

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.

Navodilo za uporabo

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

Označevanje

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