File downloaded on 2026-11-16
Source URL: https://medicines.health.europa.eu/veterinary/en/700000225711

Quinox 6 mg/g oral powder for cattle and sheep

Authorised

- Decoquinate

Product identification

Medicine name:
Quinox 6 mg/g oral powder for cattle and sheep

Active substance:
Decoquinate

Target species:
Cattle
Sheep

Route of administration:
Oral use

Product details

Active substance and strength:
Decoquinate
6.00 milligram(s) / 1.00 gram(s)

Pharmaceutical form:
Oral powder

Withdrawal period by route of administration:

Oral use:

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QP51BX04

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Cyprus

Package description:
Zipped 1 kg heat-sealed bag made of low-density polyethylene/polyamide/aluminium/polyester
5 kg heat-sealed bag made of low-density polyethylene/aluminium/polyamide

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Hybrid application – change in strength (Article 19(1)(a) of Regulation (EU) 2019/6)

Marketing authorisation holder:
HuVepharma

Marketing authorisation date:
28/07/2026

Manufacturing sites for batch release:
Huvepharma S.A.

Responsible authority:
Veterinary Services, Ministry Of Agriculture, Natural Resources And Environment

Authorisation number:
CY01030V

Date of authorisation status change:
30/06/2026

Reference member state:
Ireland

Procedure number:
IE/V/0599/001

Concerned member states:
Belgium Cyprus Czechia France Germany Greece Hungary Italy Netherlands Poland Portugal Slovakia Spain United Kingdom (Northern Ireland)

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Combined File of all Documents

This document does not exist in this language (English). You can find it in another language below.