

# 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

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**Active substance:**

Decoquinate

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**Target species:**

Cattle

Sheep

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**Route of administration:**

Oral use

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## Product details

**Active substance and strength:**

Decoquinate

6.00 milligram(s) / 1.00 gram(s)

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**Pharmaceutical form:**

Oral powder

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**Withdrawal period by route of administration:****Oral use:**

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**Cattle**

- Meat and offal. 0 day

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**Sheep**

- Meat and offal. 0 day

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QP51BX04

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Cyprus

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**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

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Hybrid application – change in strength (Article 19(1)(a) of Regulation (EU) 2019/6)

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**Marketing authorisation holder:**

HuVepharma

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**Marketing authorisation date:**

28/07/2026

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**Manufacturing sites for batch release:**

Huvepharma S.A.

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**Responsible authority:**

Veterinary Services, Ministry Of Agriculture, Natural Resources And Environment

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**Authorisation number:**

CY01030V

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**Date of authorisation status change:**

30/06/2026

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**Reference member state:**

Ireland

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**Procedure number:**

IE/V/0599/001

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**Concerned member states:**

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

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To consult adverse reactions on veterinary medicinal products please go to  
[www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Combined File of all Documents

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