Diurizone Powder

Not authorised

- Hydrochlorothiazide
- Dexamethasone

Product identification

Medicine name:
Diurizone Powder

Active substance:
Hydrochlorothiazide

Dexamethasone

Target species:
Cattle
Horse (non food-producing)

Route of administration:
Oral use

Product details

Active substance and strength:
Hydrochlorothiazide
75.00 milligram(s) / 1.00 gram(s)

Dexamethasone
0.25 milligram(s) / 1.00 gram(s)

Pharmaceutical form:
Oral powder

Withdrawal period by route of administration:

Oral use:

- **Cattle**
  - Meat and offal. 28 day
  - Milk. 7 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QC03AX01

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

Authorisation status:
Surrendered

Authorised in:
Ireland

Package description:
20g polyethylene paper sachet. Twenty sachets per cardboard box.
20g polyethylene paper sachet. Four per cardboard box.

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Vetoquinol Ireland Limited

Marketing authorisation date:
11/10/1989

Manufacturing sites for batch release:
Vetoquinol S.A.
Vetoquinol Biowet Sp. z o.o.

Responsible authority:
Health Products Regulatory Authority

Authorisation number:
VPA10983/010/001

Date of authorisation status change:
1/12/2025

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

Documents

Summary of Product Characteristics