

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:**

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Cattle

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