

DIURIZONE POUDRE

Not authorised

- Hydrochlorothiazide
- Dexamethasone

Product identification

Medicine name:

DIURIZONE POUDRE

Active substance:

Hydrochlorothiazide

Dexamethasone

Target species:

Cattle

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:

Powder for oral suspension

Withdrawal period by route of administration:

Oral use:

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Cattle

- Meat and offal. 6 day
- Milk. 3 day

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Cattle

- Meat and offal. 6 day
 - Milk. 3 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:

France

Package description:

Available only in [French](#)

Available only in [French](#)

Available only in [French](#)

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:

Vetoquinol S.A.

Marketing authorisation date:

30/06/1992

Manufacturing sites for batch release:

Vetoquinol Biowet Sp. z o.o.

Vetoquinol S.A.

Responsible authority:

French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:

FR/V/3978562 4/1992

Date of authorisation status change:

26/08/2025

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

Documents

Summary of Product Characteristics

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