

Trilorale 10 mg/ml - Oral suspension

Authorised

- Trilostane

Product identification

Medicine name:

Trilorale 10 mg/ml - Oral suspension

Active substance:

Trilostane

Target species:

Dog

Route of administration:

Oral use

Product details

Active substance and strength:

Trilostane

10.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Oral suspension

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QH02CA01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Austria , Belgium , Bulgaria , Croatia , Cyprus , Czechia , Denmark , Estonia , Finland , France , Germany , Greece , Hungary , Iceland , Ireland , Italy , Latvia , Liechtenstein , Lithuania , Luxembourg , Malta , Netherlands , Norway , Poland , Portugal , Romania , Slovakia , Slovenia , Spain , Sweden , United Kingdom (Northern Ireland)

Available in:

France

Package description:

Packaging: Bottle with 1 ml and 5 ml PP syringe (HDPE), Package_size: 1 bottle,
Content: 30 ml

Packaging: Bottle with 1 ml and 5 ml PP syringe (HDPE), Package_size: 1 bottle,
Content: 90 ml

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Hybrid application - change in pharmaceutical form (Article 19(1)(a) of Regulation (EU) 2019/6)

Marketing authorisation holder:

AXIENCE

Marketing authorisation date:

6/05/2024

Manufacturing sites for batch release:

Lelypharma B.V.

Responsible authority:

European Commission

Authorisation number:

This information is not available for this product.

Date of authorisation status change:

6/05/2024

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

Documents

Combined File of all Documents

English (PDF)

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