

Vetoryl 5 mg hard capsules for dogs

Authorised

- Trilostane

Product identification

Medicine name:

Vetoryl 5 mg hard capsules for dogs
Vetoryl 5 mg Kapsułka, twarda

Active substance:

Trilostane

Target species:

Dog

Route of administration:

Oral use

Product details

Active substance and strength:

Trilostane
5.00 milligram(s) / 1.00 Capsule

Pharmaceutical form:

Capsule, hard

Withdrawal period by route of administration:

Oral use:

-

Dog

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QH02CA01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Poland

Package description:

30 capsules, contained in three PVC-PVdc/aluminium foil blisters with 10 capsules/blister.

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:

Dechra Regulatory B.V.

Marketing authorisation date:

21/04/2021

Manufacturing sites for batch release:

Genera d.d.

Responsible authority:

Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

Authorisation number:

3089

Date of authorisation status change:

21/04/2021

Reference member state:

Ireland

Procedure number:

IE/V/0514/005

Concerned member states:

Austria Belgium Croatia Czechia Denmark Finland France Germany Greece
Hungary Italy Luxembourg Netherlands Norway Poland Portugal Slovakia
Slovenia Spain Sweden United Kingdom (Northern Ireland)

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

Documents

Package Leaflet

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

Labelling

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Summary of Product Characteristics

English (PDF)

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