

Vetoryl 20 mg hard capsules for dogs

Authorised

- Trilostane

Product identification

Medicine name:

Vetoryl 20 mg hard capsules for dogs

Active substance:

Trilostane

Target species:

Dog

Route of administration:

Oral use

Product details

Active substance and strength:

Trilostane

20.00 milligram(s) / 1.00 Capsule

Pharmaceutical form:

Capsule, hard

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QH02CA01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Slovakia

Package description:

PVC-PVdc/aluminium foil blisters in a cardboard box. Each blister contains 10 capsules. Pack size: 30 capsules

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Full application - known active substance (Article 8 of Regulation (EU) 2019/6)

Marketing authorisation holder:

Dechra Regulatory B.V.

Marketing authorisation date:

25/08/2025

Manufacturing sites for batch release:

Genera d.d.

Responsible authority:

Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:

96/007/DC/25-S

Date of authorisation status change:

25/08/2025

Reference member state:

Ireland

Procedure number:

IE/V/0514/010/DX/001

Concerned member states:

Austria Belgium Croatia Czechia Denmark Finland France Germany Greece
Hungary Italy Luxembourg Netherlands 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

Combined File of all Documents

English (PDF)

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Summary of Product Characteristics

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