

Vetoryl 20 mg hard capsules for dogs

Authorised

- Trilostane

Product identification

Medicine name:

Vetoryl 20 mg hard capsules for dogs
Vetoryl 20 mg hårda kapslar till hund

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

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:

Sweden

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:

25/02/2025

Manufacturing sites for batch release:

GENERA d.d.

Responsible authority:

Swedish Medical Products Agency

Authorisation number:

66063

Date of authorisation status change:

25/02/2025

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

Summary of Product Characteristics

English (PDF)

Published on: 13/04/2025

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Package Leaflet

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Labelling

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Combined File of all Documents

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