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Vetoryl 120 mg hard capsules for dogs

Authorised

- Trilostane

Product identification

Medicine name:
Vetoryl 120 mg hard capsules for dogs

Active substance:
Trilostane

Target species:
Dog

Route of administration:
Oral use

Product details

Active substance and strength:
Trilostane
120.00 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:
Norway

Available in:
Norway

Package description:
PVC – PVdc / aluminium foil blister packs containing 10 capsules. Each carton contains 3 blister strips.

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:
16/05/2006

Manufacturing sites for batch release:
Genera d.d.

Responsible authority:
Norwegian Medical Products Agency

Authorisation number:

05-3600

Date of authorisation status change:
22/01/2019

Reference member state:
Ireland

Procedure number:
IE/V/0514/003

Concerned member states:
Austria Belgium Czechia Denmark Finland Germany Greece Hungary Italy Luxembourg Netherlands Norway Poland Portugal Slovakia Spain Sweden United Kingdom (Northern Ireland)

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Documents

Summary of Product Characteristics

English (PDF)
Published on: 26/08/2024
Updated on: 14/03/2026
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Package Leaflet

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