# Vetoryl 60 mg hard capsules for dogs

Authorised

Trilostane

## Product identification

#### **Medicine name:**

Vetoryl 60 mg hard capsules for dogs Vetoryl 60 mg kapseli, kova

#### **Active substance:**

**Trilostane** 

### **Target species:**

Dog

#### Route of administration:

Oral use

# **Product details**

## **Active substance and strength:**

Trilostane 60.00 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: Finland
Package description: PVC-PVdc/aluminium foil blister packs containing 3 strips of 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: 19/10/2006
Manufacturing sites for batch release: GENERA d.d.
Responsible authority: Finnish Medicines Agency

Authorisation number: 21444
Date of authorisation status change: 19/10/2006
Reference member state: Ireland
Procedure number: IE/V/0514/004
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
Summary of Product Characteristics
This document does not exist in this language (English). You can find it in another language below.
Package Leaflet
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language below.

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