Vetoryl 60 mg chewable tablets for dogs

Authorised

- Trilostane

Product identification

Medicine name:
Vetoryl 60 mg chewable tablets for dogs
Vetoryl 60 mg tabletki do rozgryzania i żucia dla psów 60 mg Tabletka do rozgryzania i żucia

Active substance:
Trilostane

Target species:
Dog

Route of administration:
Oral use

Product details

Active substance and strength:
Trilostane
60.00 milligram(s) / 1.00 Tablet

Pharmaceutical form:

Chewable tablet

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:
Aluminium – Polyamide/Aluminium/PVC blister.Each blister contains 10 tablets.
Cardboard box of 1 blister.
Aluminium – Polyamide/Aluminium/PVC blister.Each blister contains 10 tablets.
Cardboard box of 3 blisters.
Aluminium – Polyamide/Aluminium/PVC blister.Each blister contains 10 tablets.
Cardboard box of 5 blisters.
Aluminium – Polyamide/Aluminium/PVC blister.Each blister contains 10 tablets.
Cardboard box of 6 blisters.
Aluminium – Polyamide/Aluminium/PVC blister.Each blister contains 10 tablets.
Cardboard box of 10 blisters.

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:
22/11/2024

Manufacturing sites for batch release:
Lelypharma B.V.

Responsible authority:
Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

Authorisation number:
3368

Date of authorisation status change:
22/11/2024

Reference member state:
Ireland

Procedure number:
IE/V/0514/008

Concerned member states:
Austria Belgium Bulgaria Croatia Cyprus Czechia Denmark Estonia Finland France Germany Greece Hungary Italy Latvia Lithuania Luxembourg Netherlands Norway Poland Portugal Romania 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

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

Labelling

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

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