Vetoryl 60 mg chewable tablets for dogs

Authorised

- Trilostane

Product identification

Medicine name:
Vetoryl 60 mg chewable tablets for dogs
Vetoryl 60 mg žvečljive tablete za pse

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

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:
Slovenia

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:
27/05/2024

Manufacturing sites for batch release:
Lelypharma B.V.

Responsible authority:
Agency For Medicinal Products And Medical Devices Of The Republic Of Slovenia

Authorisation number:
MR/V/0606/007

Date of authorisation status change:
27/05/2024

Reference member state:
Ireland

Procedure number:
IE/V/0514/008

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

English (PDF)

Published on: 25/09/2024
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Package Leaflet

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Labelling

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