Vetoryl 120 mg chewable tablets for dogs

Authorised

- Trilostane

Product identification

Medicine name:
Vetoryl 120 mg chewable tablets for dogs
Vetoryl 120 mg košļājamās tabletes suņiem

Active substance:
Trilostane

Target species:
Dog

Route of administration:
Oral use

Product details

Active substance and strength:
Trilostane
120.00 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

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Authorisation status:

Valid

Authorised in:

Latvia

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:
24/04/2025

Manufacturing sites for batch release:
Lelypharma B.V.

Responsible authority:
Food And Veterinary Service

Authorisation number:
V/SRP/25/0028

Date of authorisation status change:
24/04/2025

Reference member state:
Ireland

Procedure number:
IE/V/0514/009

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

Combined File of all Documents

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