Source URL: https://medicines.health.europa.eu/veterinary/en/700000130151

Vetoryl 120 mg chewable tablets for dogs

Authorised

Trilostane

Product identification

Medicine name:

Vetoryl 120 mg chewable tablets for dogs Vetoryl 120 mg chewable tablets 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 Tablet

Pharmaceutical form:

Chewable tablet

Withdrawal period by route of administration:

Oral use:

Dog

Anatomical therapeutic chemical veterinary (ATCvet) codes:

OH02CA01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Ireland

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: 28/06/2024 |
| Manufacturing sites for batch release: Lelypharma B.V. |
| Responsible authority: Health Products Regulatory Authority |
| Authorisation number: VPA22622/023/009 |
| Date of authorisation status change: 28/06/2024 |
| 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 |

Summary of Product Characteristics

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