Vetoryl 30 mg chewable tablets for dogs

Authorised

Trilostane

Product identification

Medicine name:

Vetoryl 30 mg chewable tablets for dogs Vetoryl 30 mg chewable tablets for dogs

Active substance:

Trilostane

Target species:

Dog

Route of administration:

Oral use

Product details

Active substance and strength:

Trilostane

30.00 milligram(s) / 1.00 Tablet

Pharmaceutical form:

Chewable tablet

Withdrawal period by route of administration:

Oral use:

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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: VPA22622/023/007
Date of authorisation status change: 28/06/2024
Reference member state: Ireland
Procedure number: IE/V/0514/007
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

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