Source URL: https://medicines.health.europa.eu/veterinary/en/700000179563

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:

OH02CA01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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

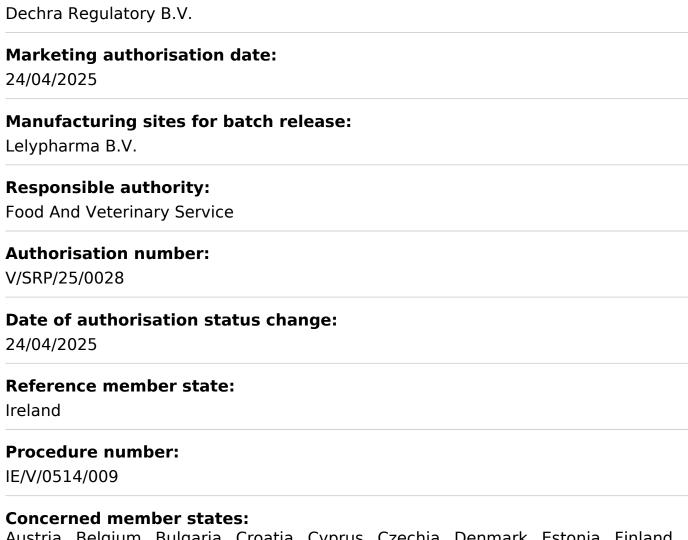
Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Full application - known active substance (Article 8 of Regulation (EU) 2019/6)

Marketing authorisation holder:



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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