Vetoryl 30 mg hard capsules

Authorised

- Trilostane

Product identification

Medicine name:
Vetoryl 30 mg hard capsules
Vetoryl 30 mg trde kapsule za pse

Active substance:
Trilostane

Target species:
Dog

Route of administration:
Oral use

Product details

Active substance and strength:
Trilostane
30.00 milligram(s) / 1.00 Capsule

Pharmaceutical form:
Capsule, hard

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:
PVC-PVdc / aluminium foil blister packs containing 10 capsules. Each carton contains 3 blister strips.

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Dechra Regulatory B.V.

Marketing authorisation date:
2/02/2018

Manufacturing sites for batch release:
Genera d.d.

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

Authorisation number:
MR/V/0606/002

Date of authorisation status change:
2/02/2018

Reference member state:
Ireland

Procedure number:
IE/V/0514/002

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: 23/08/2024
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Package Leaflet

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Labelling

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